TITLE OF THE INVENTION

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METHOD FOR LOW TEMPERATURE BONDING AND BONDED STRUCTURE

CROSS REFERENCE TO RELATED APPLICATIONS

This application is a continuation of, and claims the benefit of priority under 35 U.S.C. § 120 from U.S. Serial No. 14/754,111, filed June 29, 2015, which is a division of U.S. Serial No. 14/197,070, filed March 4, 2014, which is a division of U.S. Serial No. 13/341,273, filed December 30, 2011, which is a continuation of U.S. Serial No. 12/954,740, filed November 26, 2010, now Patent No. 8,153,505, which is a continuation of U.S. Serial No. 12/720,368 filed March 9, 2010, now Patent No. 7,871,898, which is a continuation of U.S. Serial No. 11/980,664 filed October 31, 2007, now Patent No. 7,807,549, which is a continuation of U.S. Serial No. 10/913,441 filed August 9, 2004, now Patent No. 7,387,944, which is a continuation of U.S. Serial No. 09/505,283, filed February 16, 2000, now Patent No. 6,902,987, the entire contents of each of which are incorporated herein by reference.

BACKGROUND OF THE INVENTION

1. Field of the Invention

The present invention relates to bonding of materials at room temperature and, in particular, to bonding of processed semiconductor materials, such as integrated circuit or device substrates, having activated surfaces to achieve high bonding strength adequate for subsequent fabrication and/or a desired application.

2. Background of the Invention

Direct room temperature bonding generally produces weak van der Waals or hydrogen bonding. Annealing is typically required to convert the weak bond to a stronger chemical bond such as a covalent bond. Other wafer bonding techniques including anodic and fusion typically require the application of voltage, pressure and/or annealing at elevated temperature to achieve a sufficient bond strength for subsequent fabrication and/or the desired application. The need to apply voltage, pressure or heat has significantly limited wafer bonding applications because these parameters can damage the materials being wafer bonded, give rise to internal stress and introduce undesirable changes in the devices or materials being bonded. Achieving a strong bond at low temperatures is also critical for

bonding of thermally mismatched or thermally sensitive wafers including processed device wafers.

Ultra high vacuum (UHV) bonding is one of the approaches to achieve a low or room temperature strong bond. However, the bonding wafers still have to be pre-annealed at high temperatures, for instance >600°C for silicon and 500°C for GaAs, before cooling down to low or room temperature for bonding. Furthermore, the UHV approach does not generally work on commonly used materials, for example, in SiO₂. It is further also expensive and inefficient.

Adhesive layers can also be used to bond device wafers to a variety of substrates and to transfer device layers at low temperatures. However, thermal and chemical instability, interface bubbles, stress and the inhomogeneous nature of adhesive layers prevent its wide application. It is thus highly desirable to achieve a strong bond at room temperature by bonding wafers in ambient without any adhesive, external pressure or applied electric field.

Low vacuum bonding has been explored as a more convenient alternative to UHV bonding but a bonding energy comparable to the bulk silicon fracture energy using bonded bare silicon wafer pairs has only be achieved after annealing at $\sim 150^{\circ}$ C. For oxide covered silicon wafer pairs annealing at $\sim 300^{\circ}$ C is required to obtain a high bond energy. It has not been possible to obtain high bonding energies in bonded material using low vacuum bonding at room temperature.

A gas plasma treatment prior to bonding in ambient is known to enhance the bonding energy of bonded silicon pairs at low or room temperature. See, for example, G.L. Sun, Q.-Y. Tong, et al., J. de Physique, 49(C4), 79 (1988); G.G. Goetz, Proc. of 1st Symp. on Semicond. Wafer Bonding: Science, Technol. and Applications, The Electrochem. Soc., 92-7, 65 (1992); S. Farrens et al., J. Electroch. Soc., 142,3950 (1995) and Amirffeiz et al, Abstracts of 5th Symp. on Semi. Wafer Bonding: Science, Tech. and Appl., The Electrochemical Society, 99-2, Abstract No. 963 (1999). Although these treatments have increased the bond energy obtainable at low or room temperature, they have only been demonstrated with planar silicon wafers or with silicon wafers using a plasma process that results in oxide being grown on the wafers during the plasma process. Moreover, these treatments have only been used to increase the bond energy by charging or damaging the surface. Furthermore, these treatments have not been used or shown to be applicable to deposited dielectrics or other materials.

Obtaining low or room temperature bonding with a method that is not only applicable to planar silicon and grown oxide surfaces but further to deposited materials and non-planar

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surfaces with planarized deposited materials will allow generic materials, including processed semiconductor wafers, to be bonded with minimal damage for manufacturing purposes. Such a method based on etching and chemical bonding is described herein.

SUMMARY OF THE INVENTION

It is an object of the invention to provide a method for bonding materials at low or room temperature.

It is another object of the invention to bond materials by cleaning and activating the bonding surfaces to promote chemical bond formation at about room temperature.

It is a further object of the invention to provide a bonding method to bond any solid state material such as processed device or integrated circuit wafers or thermally sensitive or mis-matched materials at or about room temperature.

It is further object of the invention to provide a bonding method to bond processed device or integrated circuit wafers of different types of devices or different technologies, and transfer a layer of devices or circuits at or about room temperature.

It is another object of the invention to enable a direct wafer bonding method that does not require annealing to achieve a required bond strength.

It is a further object of the invention to provide a method whereby diverse materials including those with non-planar surfaces and deposited materials can be planarized and bonded.

These and other objects are achieved by a method of bonding having steps of forming first and second bonding surfaces, etching the first and second bonding surfaces, and

bonding together at room temperature the first and second bonding surfaces after said etching step. The etching may include etching the first and second bonding surfaces such that respective surface roughnesses of the first and second bonding surfaces after said etching are substantially the same as respective surface roughnesses before said etching. The surface roughness may be in a range of 0.1 to 3.0 nm.

The bonding surfaces may be the surface of a deposited insulating material, such as silicon oxide, silicon nitride or a dielectric polymer. The bonding surface may also be the surface of a silicon wafer. Silicon wafers, using either the surface of the wafer or a deposited material on the wafer, may be bonded together. The wafers may have devices or integrated circuits formed therein. The devices and circuits in the wafers bonded together may be

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interconnected. The wafers may have a non-planar surface or an irregular surface topology upon which a material is deposited to form the bonding surfaces.

Forming at least one of the bonding surfaces may include depositing a polishable material on a non-planar surface. Depositing said polishable material may include depositing one of silicon oxide, silicon nitride or a dielectric polymer. The bonding surfaces may be polished using a method such as chemical-mechanical polishing. The surfaces may also be etched prior to the polishing.

The etching step may also include activating the first and second bonding surfaces and forming selected bonding groups on the first and second bonding surfaces. Bonding groups may also be formed capable of forming chemical bonds at approximately room temperature, and chemical bonds may be formed between the bonding surfaces allowing bonded groups to diffuse or dissociate away from an interface of the bonding surfaces. The chemical bonds can increase the bonding strength between the bonding surfaces by diffusing or dissociating away said bonding groups.

After said etching step, the bonding surfaces may be immersed in a solution to form bonding surfaces terminated with desired species. The species may comprise at least one of a silanol group, an NH₂ group, a fluorine group and an HF group. Also, a monolayer of one of a desired atom and a desired molecule may be formed on the bonding surface. Terminating the surface may include rinsing said bonding materials in an ammonia-based solution after said slightly etching. The ammonia-based solution may be ammonium hydroxide or ammonium fluoride.

The method may also include exposing the bonding surfaces to one of an oxygen, argon, NH₃ and CF₄ RIE plasma process. Silicon dioxide may be deposited as to form the bonding surfaces, and etched using the RIE process.

The etching process may create a defective or damaged zone proximate to the bonding surfaces. The defective or damaged zone can facilitate the removal of bonding by-products through diffusion or dissociation.

The method may also include steps of forming first and second bonding surfaces, etching the bonding surfaces, terminating the bonding surfaces with a species allowing formation of chemical bonds at about room temperature, and bonding the bonding surfaces at about room temperature, or may include steps of forming the bonding surfaces each having a surface roughness in a range of 0.1 to 3 nm, removing material from the bonding surfaces while maintaining said surface roughness, and directly bonding the bonding surfaces at room

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temperature with a bonding strength of at least 500 mJ/m², at least 1000 mJ/m², or at least 2000 mJ/m².

The objects of the invention may also be achieved by a bonded device having a first material having a first etched bonding surface, and a second material having a second etched bonding surface directly bonded to the first bonding surface at room temperature having a bonding strength of at least 500 to 2000 mJ/m². The bonding surfaces may be being activated and terminated with a desired bonding species, and the desired species may include a monolayer of one of a desired atom and a desired molecule on said bonding surface or at least one of a silanol group, an NH₂ group, a fluorine group and an HF group. The bonding surfaces may each have a defective region located proximate to said first and second bonding surfaces, respectively.

The first material may include a surface of a first semiconductor wafer having devices formed therein, and the second material may include a surface of a second semiconductor wafer having devices formed therein. Devices in the wafers may be interconnected, and the wafers may be of different technologies. The wafers may also have an integrated circuit formed therein, and devices or circuits in the wafers may be interconnected.

One of said first and second wafers may be a device region after removing a substantial portion of a substrate of said one of said first and second wafers. The wafers may have an irregular surface topology.

The first material may include a first wafer containing electrical devices and having a first non-planar surface, and the first bonding surface may include a polished and etched deposited oxide layer on said first non-planar surface. The second material may include a second wafer containing electrical devices and having a second non-planar surface, and the second bonding surface may include a polished, planarized and slightly etched deposited oxide layer on the second non-planar surface.

The first material may include a first wafer containing electrical devices and having a first surface with irregular topology, and the first bonding surface may include a polished, planarized and slightly etched deposited oxide layer on the first surface. The second material may include a second wafer containing electrical devices and having a second surface with irregular topology, and the second bonding surface may include a polished, planarized and slightly etched deposited oxide layer on the second surface.

The bonded device according to the invention may also include a first material having a first etched and activated bonding surface terminated with a first desired bonding species,

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and a second material having a second etched and activated bonding surface terminated with a second desired bonding species bonded to the first bonding surface at room temperature.

BRIEF DESCRIPTION OF THE DRAWINGS

- A more complete appreciation of the invention and many of the attendant advantages thereof are readily obtained as the same become better understood by reference to the following detailed description when considered in connection with the accompanying drawings, wherein:
 - FIG. 1 is a flow chart of the method according to the invention;
 - FIG. 2 is a flow chart of an example of the method according to the invention;
 - FIGS. 3A-3E are diagrams illustrating a first embodiment of a method according to the invention:
 - FIGS. 4A-4E are diagrams illustrating bonding according to the invention using silicon oxide;
- FIGS. 5A-5E are diagrams illustrating bonding according to the invention using silicon;
 - FIGS. 6A and 6B are graphs of room temperature bonding energy versus storage time;
 - FIG. 7 is a diagram of a bonding fixture used in the invention; and
 - FIG. 8 is a fluorine concentration profile by SIMS (Secondary Ion Mass
- Spectroscopy) near the bonding interface of deposited oxide covered silicon wafers that were very slight etched by diluted HF before bonding.

DESCRIPTION OF THE PREFERRED EMBODIMENTS

Referring to FIGS. 1 and 3A-3E, a first embodiment of the method according to the invention will be described. Wafer 30, preferably a processed semiconductor device wafer and more preferably a processed silicon device wafer, contains a device layer 31 with processed devices. Device layer 31 may contain a number of layers and include surface regions of wafer 30. The surface topography of layer 31 is typically nonplanar. Layer 31 may also represent a processed integrated circuit containing any number of layers such as active devices, interconnection, insulation, etc.

The integrated circuit may be fully processed, or partially processed where the remaining processing is performed after the bonding process. The processing after the

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bonding may include full or partial substrate removal or via formation between the bonded wafers for interconnection.

On layer 31 a bonding layer 32 is formed (step 1, FIG. 1). Bonding layer 32 may be any solid state material or mixed materials which can be deposited or formed at low temperatures and can be polished to a sufficiently smooth surface. Layer 32 may be an insulator, such as SiO₂, silicon nitride, amorphous silicon formed using chemical vapor deposition (CVD) or plasma-enhanced CVD (PECVD), sputtering or by evaporation. Other materials such as polymers, semiconductors or sintered materials may also be used. Layer 32 should have thickness greater than the surface topography of layer 31.

The surface 33 of layer 32 is planarized and smoothed, as shown in step 2 of FIG. 1 and in FIG. 3B. It is noted that the roughness/planarity of surface 33 is exaggerated in FIG. 3A for illustrative purposes. This step may be accomplished using chemical-mechanical polishing. Surface 33 is preferably polished to a roughness of about no more than about 3 nm and preferably no more than about 0.1 nm and be substantially planar. The surface roughness values are typically given as root-mean square (RMS) values. Also, the surface roughness may be given as mean values which are nearly the same as the RMS values. After polishing surface 33 is cleaned and dried to remove any residue from the polishing step. Polished surface 33 is preferably then rinsed with a solution.

The bonding surface may also be etched prior to polishing to improve the planarity and/or surface roughness. The etching can be effective to remove high spots on the bonding surface by selective etching of the high spots using, for example, standard photolithographic techniques. For example, a layer of silicon nitride can be embedded within a silicon dioxide bonding layer 32 that can serve as an etch stop when using a solution containing HF. The etch stop material may be used to improve uniformity, reproducibility, and manufacturability.

FIG. 3B illustrates layer 32 having upper surface 34 after the polishing/planarization and cleaning steps. Surface 34 then undergoes an activation process (step 3, FIG. 1). This activation process is an etching process and preferably a very slight etch (VSE) process. The term VSE means that the root-mean-square micro-roughness (RMS) of the very slightly etched surface remains at approximately the unetched value, typically < 0.5 nm and preferably in the range of 0.1 nm to 3 nm. The optimum amount of material removed depends upon the material and the method used for removal. Typical amounts removed vary from Angstroms to a few nanometers. It is also possible to remove more material. VSE also includes the breaking of bonds on the treated surfaces and can occur without significant

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removal of material. The VSE is distinct from simple modification of the surface by, for example, charging the surface with electronic charge or damaging the surface layer. In a first example of the method according to the invention, the VSE process consists of a gas or mixed gas (such as oxygen, argon, nitrogen, CF₄, NH₃) plasma process at a specified power level for a specified time (FIG. 3C). The power and duration of the plasma process will vary depending upon the materials used to obtain the desired bond energy. Examples are given below, but in general, the power and duration will be determined empirically.

The plasma process may be conducted in different modes. Both reactive ion etch (RIE) and plasma modes may be used, as well as an inductively-coupled plasma mode (ICP). Sputtering may also be used. Data and examples are given below in both the RIE and plasma modes.

The VSE process etches the surface very slightly via physical sputtering and/or chemical reaction and preferably is controlled to not degrade the surface roughness of the bonding surfaces. The surface roughness may even be improved depending upon the VSE and materials etched. Almost any gas or gas mixture that will not etch surface 34 excessively can be used for the room temperature bonding method according to the invention.

The VSE serves to clean the surface and break bonds of the oxide on the wafer surface. The VSE process can thus enhance the surface activation significantly. A desired bonding species can be used to terminated on surface 34 during the VSE by proper design of the VSE. Alternatively, a post-VSE treatment that activates and terminates the surface with a desired terminating species during the post-VSE process may be used.

The desired species further preferably forms a temporary bond to the surface 34 atomic layer, effectively terminating the atomic layer, until a subsequent time that this surface can be brought together with a surface terminated by the same or another bonding species 36 as shown in FIG. 3D. Desired species on the surfaces will further preferably react with each other when they are in sufficiently close proximity allowing chemical bonding between surfaces 34 and 36 at low or room temperature that is enhanced by diffusion or dissociation and diffusion of the reacted desired species away from the bonding interface.

The post-VSE process preferably consists of immersion in a solution containing a selected chemical to generate surface reactions that result in terminating the bonding surface 34 with desired species. The immersion is preferably performed immediately after the VSE process. The post-VSE process may be performed in the same apparatus in which the VSE process is conducted. This is done most readily if both VSE and post-VSE processes are

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either dry, i.e, plasma, RIE, ICP, sputtering, etc, or wet, i.e., solution immersion. A desired species preferably consists of a monolayer or a few monolayers of atoms or molecules.

The post-VSE process may also consist of a plasma, RIE, or other dry process whereby appropriate gas chemistries are introduced to result in termination of the surface with the desired species. The post-VSE process may also be a second VSE process. The termination process may also include a cleaning process where surface contaminants are removed without VSE. In this case, a post-cleaning process similar to the post-VSE processes described above then results in a desired surface termination.

The post-VSE or post-cleaning process may or may not be needed to terminate surfaces with desired species if the activated surface bonds by the cleaning or VSE process are subsequently sufficiently weakly surface reconstructed and can remain sufficiently clean before bonding such that subsequent bonding with a similar surface can form a chemical bond.

The wafers are optionally rinsed then dried. Two wafers are bonded by aligning them (if necessary) and bringing them together to form a bonding interface. As shown in FIG. 3D, a second wafer 35 has been processed in the manner shown in FIG. 3C to prepare bonding surface 36. The two wafers are brought together by, for example, commercially available wafer bonding equipment (not shown) to initiate bonding interface 37 (FIG. 3E).

A spontaneous bond then typically occurs at some location in the bonding interface and propagates across the wafer. As the initial bond begins to propagate, a chemical reaction such as polymerization that results in chemical bonds takes place between species used to terminate surfaces 34 and 36 when the surfaces are in sufficient proximity. The bonding energy is defined as the specific surface energy of one of the separated surfaces at the bonding interface that is partially debonded by inserting a wedge. The by-products of the reaction then diffuse away from the bonding interface to the wafer edge or are absorbed by the wafers, typically in the surrounding materials. The by-products may also be converted to other by-products that diffuse away or are absorbed by the wafers. The amount of covalent and/or ionic bonding may be increased by removal of converted species resulting in further increase in bond strength.

FIGS. 4A-4E show surface conditions and the bonding propagation to form covalent bonds in a the case of a planar Si wafer covered with silicon oxide. On Si wafer 40 an SiO₂ layer 41 is formed, which has been polished and planarized. Surface 42 of layer 41 is subjected to the VSE process to produce an activated surface (FIG. 4A). On a second wafer

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44 a second SiO₂ layer 45 is formed, and surface 46 is subjected to a VSE process to activate surface 46 (FIG. 4B). Desired species are terminated on surface 46 and are shown as lines 43 in FIG. 4C. Either or both of a VSE and post-VSE processes are used to properly terminate surface 46. While not shown, surface 42 may also be terminated using a post-VSE process. Wafer 44 is brought together with wafer 40 (FIG. 4D) and bonds 46 begin to form. The bonding propagates and by-products are removed (indicated as arrows 47) and chemical bonds (such as covalent) are formed, as shown in FIG. 4E.

The bonding immediately after the RIE process may use a special bonding fixture allowing immediate in situ bonding of the etched wafers. A diagram of the fixture is shown in FIG. 7. In plasma chamber 75 are two wafers to be bonded 70 disposed on RF electrodes 76 and 77. A plasma is formed in zone 79 by the application of RF power to the electrodes via moveable vacuum RF power feedthrough 74 and by the introduction of an appropriate gas or gas mixture through gas feedthrough 73. Element 71 is a vacuum feedthrough for mechanical actuator (not shown) to retract retractable spacer 72. Chamber 75 is pumped down to a desired vacuum level via pumps (not shown) and chamber inlet 78. In the case where a post-VSE process or post cleaning process is also a dry process, as discussed above, the VSE and post-VSE or post-cleaning may be conducted in chamber 75.

After the plasma treatment to conduct the VSE process, the mechanical spacers 72 are retracted by the mechanical actuator and the wafers 70 are moved into contact with to begin the bonding process. The bonded wafers are then moved from the chamber into ambient or into another vacuum chamber (not shown) and stored for a desired period to allow the bonding to propagate by a wafer handling system (not shown).

The materials of the bonding layers preferably have an open structure so that the byproducts of the polymerization reaction can be easily removed. The bonding species on the
opposing bonding surfaces must be able to react at room temperature to form a strong or
chemical bond. The bond energy is sufficiently high to virtually eliminate slippage between
wafers after subsequent heat treatments associated with a subsequent processing or operation
when wafers have different thermal expansion coefficients. Lack of slippage is manifest by a
lack of wafer bowing upon inspection after the subsequent processing or operation.

In order to achieve the high bonding energies, it is preferable for at least one of the wafers to be as thin as possible because a thin wafer allows compliance to accommodate a lack of perfect surface planarization and smoothness. Thinning to thickness of about 10 mils to 10 microns is effective.

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The bonded wafers are preferably stored at ambient or at low or room temperature after bonding to allow removal of species or converted species for a specified period of time depending upon the materials and species used. Twenty four hours is usually preferable. The storage time is dependent upon the type of plasma process used. Chemical bonds may be obtained more quickly, in a matter of minutes, when certain plasma processes such as an Ar plasma are used. For example, 585 mJ/m² bonds were obtained in immediately after bonding and over 800 mJ/m² were observed after 8 hours for deposited oxides etched by an Ar plasma followed by NH₄OH dip.

Annealing the bonded wafers during bonding may increase the bonding strength. The annealing temperature should be below 200EC and may be typically in the range of 75-100EC. Storing the bonded wafers under vacuum may facilitate the removal of residual gasses from the bonding surfaces, but is not always necessary.

All of the processes above may be carried out at or near room temperature. The wafers are bonded with sufficient strength to allow subsequent processing operations (lapping, polishing, substrate removal, chemical etching, lithography, masking, etc.). Bonding energies of approximately 500-2000 mJ/m² or more can be achieved (see FIG. 6A).

At this point (FIG. 3E) it is possible to remove a part or all of the substrate of wafer 35 by, for instance, lapping and etch back. The layer of devices of wafer 35 is thus transferred onto wafer 30. The devices from the two layers may be interconnected. Additional device or circuit layers may be bonded and interconnected to form a multilayer structure. Different types of wafers, devices or circuits may be bonded, as well as different technologies (i.e. CMOS and bipolar or III-V HBT and Si CMOS). Other elements or materials such as thermal spreaders, surrogate substrates, antennas, wiring layers, a preformed multi-layer interconnects, etc. may be bonded to produce different types of circuits or systems, as desired.

In an example, shown in FIG. 2, PECVD SiO₂ is deposited on a Si wafer containing devices. Surface 34, after the plasma (such as argon, oxygen or CF₄) treatment, is mainly terminated by Si-OH groups due to the availability of moisture in the plasma system and in air. After the plasma treatment, the wafers are immediately immersed in solution such as ammonium hydroxide (NH₄OH), NH₄F or HF for a period such as between 10 and 120 seconds. After immersing the wafers in the NH₄OH solution, many Si-OH groups are replaced by Si-NH2 groups according to the following substitution reaction:

$$2Si-OH + 2NH_4OH \rightarrow 2Si-NH_2 + 4HOH \tag{1}$$

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Alternatively, many Si-F groups are terminating on the PECVD SiO₂ surface after an NH₄F or HF immersion.

The hydrogen bonded Si-NH2:Si-OH groups or Si-NH2:Si-NH2 groups across the bonding surfaces can polymerize at room temperature in forming Si-O-Si or Si-N-N-Si (or Si-N-Si) covalent bonds:

$$Si-NH_2 + Si-OH \rightarrow Si-O-Si + NH_3$$
 (2)

$$Si-NH_2 + Si-NH_2 \rightarrow Si-N-N-Si + 2H_2$$
 (3)

Alternatively, the HF or NH₄F dipped oxide surfaces are terminated by Si-F groups in addition to Si-OH groups. Since HF or NH₄F solution etches silicon oxide strongly, their concentrations must be controlled to an adequately low level, and the immersion time must be sufficiently short. This is an example of a post-VSE process being a second VSE process. The covalent bonds across the bonding interface are formed due to the polymerization reaction between hydrogen bonded Si-HF or Si-OH groups:

$$Si-HF + Si-HF \rightarrow Si-F-F-Si + H_2$$
 (4)

$$Si-F + Si-OH \rightarrow Si-O-Si + HF$$
 (5)

FIG. 8 shows the fluorine concentration profile of bonded thermal oxide covered silicon wafers that were dipped in 0.05% HF before room temperature bonding. A fluorine concentration peak is clearly seen at the bonding interface. This provides evidence of the chemical process described above where the desired species are located at the bonding interface.

Since reaction (2) is reversible only at relatively high temperatures of \sim 500EC, the formed siloxane bonds should not be attacked by NH₃ at lower temperatures. It is known that H₂ molecules are small and diffuse about 50 times quicker than water molecules in oxide. The existence of a damaged layer near the surface of an adequate thickness i.e. a few nm, will facilitate the diffusion or dissolution of NH₃, and HF and hydrogen in reactions (2), (3), (4) and/or (5) in this layer and enhancement of the chemical bond. The three reactions result in a higher bonding energy of SiO₂/SiO₂ bonded pairs at room temperature after a period of storage time to allow NH₃ or H₂ to diffuse away.

In the example of FIG. 2, the plasma treatment may create a damaged or defective area in the oxide layer near the bonding surface. The zone extends for a few monolayers. The damaged or defective area aids in the removal of bonding by-products. Efficient

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removal of the bonding by-products improves the bonding strength since the by-products can interfere with the bonding process by preventing high-strength bond from forming.

Many different surfaces of materials may be smoothed and/or planarized, followed by a cleaning process, to prepare for bonding according to the invention. These materials can be room temperature bonded by mating surfaces with sufficient planarity, surface smoothness, and passivation that includes cleaning, and/or VSE, activation and termination. Amorphous and sintered materials, non-planar integrated circuits, and silicon wafers are examples of such materials. Single crystalline semiconductor or insulating surfaces, such as SiO₂ or Si surfaces, can also be provided with the desired surface roughness, planarity and cleanliness. Keeping the surfaces in high or ultra-high vacuum simplifies obtaining surfaces sufficiently free of contamination and atomic reconstruction to achieve the strong bonding according to the invention. Other semiconductor or insulator materials such as InP, GaAs, SiC, sapphire, etc., may also be used. Also, since PECVD SiO₂ may be deposited on many types of materials at low temperatures, many different combinations of materials may be bonded according to the invention at room temperature. Other materials may also be deposited as long as appropriate processes and chemical reactions are available for the VSE, surface activation, and termination.

For example, the method may also be used with silicon nitride as the bonding material. Silicon nitride may be bonded to silicon nitride, or to silicon dioxide and silicon. Silicon oxide may also be bonded to silicon. Other types of dielectric materials may be bonded together including aluminum nitride and diamond-like carbon.

The method may be applied to planar wafers having no devices or circuits and one wafer with devices and circuits. The planar wafer may be coated with a bonding layer, such as PECVD oxide or amorphous silicon, and then processed as described above to bond the two wafers. The planar wafer may not need to be coated with a bonding layer if it has sufficient smoothness and planarity and the proper bonding material.

As can be appreciated, the bonding process may be repeated with any number of wafers, materials or functional elements. For example, two device or IC wafers may be joined, followed by removing one of the exposed substrates to transfer a layer or more of devices, or just the active regions of an IC.

The bonding according to the invention may be applied to joining different types of materials. For example, a silicon wafer can be bonded to another silicon wafer, or bond to an oxidized silicon wafer. The bare silicon wafer and the oxide covered wafer are immersed in

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HF, NH₄F and/or NH₄OH and bonded after drying. The time for the immersion should be less than about twenty minutes for the silicon wafer covered with the thin oxide since the NH ₄OH solution etches silicon oxide. Since HF and NH₄F etches oxides strongly, very diluted solutions, preferably in 0.01-0.2% range should be used for dipping of the silicon wafers.

After drying the silicon wafer and the oxide-covered wafer are bonded in ambient at room temperature. Reactions (2), (3), (4) and/or (5) take place at the bonding interface between the two wafers. The plasma-treated wafers may also be immersed in deionized water instead of the NH₄OH solution.

The silicon bonding may be conducted with a bare silicon wafer, i.e. having a native oxide or a silicon wafer having an oxide layer formed on its surface as described above. During the oxygen plasma treatment, the native oxide which if formed on the bare silicon wafer is sputter etched, and the oxide layer formed on the silicon surface is etched. The final surface is an activated (native or formed) oxide. When rinsed in deionized water, the activated oxide surface is mainly terminated with Si-OH groups. Since oxide growth in oxygen plasma has been found to have less water than in normal native oxide layers, the water from the original bonding bridge and generated by the following polymerization reaction (6) can be absorbed into the plasma oxide readily.

$$Si-OH + Si-OH \rightarrow Si-O-Si + H_2O$$
 (6)

FIGS. 5A-5E illustrate bonding two silicon wafers. Wafers 50 and 52 have respective surfaces 51 and 53 with native oxides (not shown) subjected to a VSE process (FIGS. 5A and 5B). Surface 53 is in FIG. 5C is shown terminated with a desired species 54. The two wafers are brought together and bonds 55 begin to form (FIG. 5D). The bonding propagates and bonding by-products, in this case H₂ gas, are removed. The by-products being removed are shown as arrows 56 in FIG. 5E.

In addition to removal of the water from the bonding interface by dissolving into the plasma activated oxide of the oxidized silicon wafer, the water can also diffuse through the thin oxide layer on the bare silicon wafer to react with silicon. As the silicon surface underneath the oxide has a damaged or defective zone, extending for a few monolayers, the water molecules that diffuse through the oxide layer and reach the damaged or defective zone can be converted to hydrogen at room temperature and be removed readily:

$$Si + 2H_2O \rightarrow SiO_2 + 2H_2 \tag{7}$$

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The reverse reaction of (6) is thus avoided and the room temperature bonding energy increases enormously due to the formation of covalent Si-O-Si bonds.

If a relatively thick (~5 nm) oxide layer is formed, it will take a long period of time for the water molecules to diffuse through this thick layer. On the other hand, if after the plasma treatment a thin oxide layer is left or a too narrow defective zone is formed, water that can reach the silicon surface may not react sufficiently with the silicon and convert to hydrogen. In both cases the bonding energy enhancement will be limited. The preferred oxygen plasma treatment thus leaves a minimum plasma oxide thickness (e.g., about 0.1-1.0 nm) and a reasonably thick defective zone (e.g., about 0.1-0.3 nm) on the silicon surface.

In a second embodiment, the VSE process uses wet chemicals. For example, an InP wafer having a deposited silicon oxide layer, as in the first embodiment, and a device layer are bonded to a AlN substrate having a deposited oxide layer. After smoothing and planarizing the InP wafer bonding surface and the AlN wafer bonding surface, both wafers are cleaned in an standard RCA cleaning solution. The wafers are very slightly etched using a dilute HF aqueous solution with an HF concentration preferably in the range of 0.01 to 0.2%. About a few tenths of a nm is removed and the surface smoothness is not degraded as determined by AFM (atomic force microscope) measurements. Without deionized water rinse, the wafers are spin dried and bonded in ambient air at room temperature. The resulting bonding energy has been measured to reach ~700 mJ/m² after storage in air. After annealing this bonded pair at 75°C the bonding energy of 1500 mJ/m² was obtained. The bonding energy has been measured to reach silicon bulk fracture energy (about 2500 mJ/m²) after annealing at 100°C. If the wafers are rinsed with deionized water after the HF dip, the bonding energy at 100°C is reduced to 200 mJ/m², that is about one tenth of that obtained without the rinse. This illustrates the preference of F to OH as a terminating species.

In a third embodiment the VSE process consists of 0.1% HF etching followed by 5 min dip in 0.02% HN₄F solution of thermally oxidized silicon wafers at room temperature after a standard cleaning process. Without rinsing in deionized water, the wafers are bonded after spin drying at room temperature. The bonding energy of the bonded pairs reaches ~1700 mJ/m² after 100°C annealing. If the wafers are rinsed in de-ionized water after the HF etching before bonding, the bonding energy of bonded pairs is only 400 mJ/m², again illustrating the preference of F to OH as a terminating species. Dilute NH₄F is used in the VSE process to etch silicon oxide covered wafers in a fourth embodiment. The concentration

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of the NH₄F should be below 0.02% to obtain the desired bonding. The bonding energy of $\sim 600 \text{ mJ/m}^2$ can be achieved at room temperature after storage.

A fifth embodiment of the invention is used to bond Si surfaces having a native oxide of about 1 nm in thickness. In the fifth embodiment, after cleaning the Si surface by a standard RCA1 cleaning process, a VSE process using 5 min etching in 70% HNO₃ + diluted HF (preferably 0.01 to 0.02%) is performed. Wafers are pulled out of the solution vertically with a basically hydrophobic surface. Without rinsing in water, the wafers were bonded at room temperature in air. In this process covalent bonding occurs at room temperature with measured bonding energies typically about 600 mJ/m². This bonding energy is significantly increased to 1300 mJ/m² after annealing at 75°C and reaches the fracture energy of bulk silicon (about 2500 mJ/m²) at a temperature of 100°C.

Instead of 70% HNO₃, diluted HNO₃ with water can be used in the solution to achieve similar results. According to AMF measurements and high resolution transmission electron microscopy measurement results, the silicon is etched in the dilute HNO₃ VSE process at a rate of 0.1-0.15 nm/min and a new thick oxide 2.5-3.5 nm in thickness is formed.

As further embodiments, the VSE process may consist of a dry etch that has chemical and/or physical components. For a bare Si surface, chemical etching may result from SF₄/H₂ gas mixture while physical etching may result from Ar etch. For a silicon oxide surface, chemical etching may use CF₄ while physical etching may use oxygen or argon gas. It is also possible to use a thermally stable polymer material for the bonding materials and bond two polymer surfaces together. Examples are polyimides or spin-on materials.

The mechanisms governing the increased bond energy at low or room temperature are similar. A very slight etching (VSE) of the bonding wafers by plasma to clean and activate the surfaces, and improve removal of by-products of interface polymerization to prevent the undesirable reverse reaction and rinse in appropriate solution to terminate the surface with desired species to facilitate room temperature covalent bonding. The oxide covered wafer bonding case is similar except that a different surface termination is preferred. In bare silicon wafer bonding, the highly reactive surface layers of oxide and silicon to allow water adsorption and conversion to hydrogen should be formed. The highly reactive layers can be a plasma thin oxide layer and a damaged silicon surface layer. The oxide on the silicon wafer will also have some damage. Not only O_2 plasma but also plasma of other gases (such as Ar, CF_4) are adequate. Because during and after VSE the silicon surface is readily to react with

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moisture to form an oxide layer, and the underlying damaged silicon layer is created by VSE. Since the VSE and by-products removal methods are rather general in nature, this approach can be implemented by many means and apply to many materials.

5 EXAMPLE 1

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In a first example, three inch <100>, 1-10 ohm-cm, boron doped silicon wafers were used. PECVD oxide was deposited on some of the silicon wafers. For comparison, thermal oxidized silicon wafers were also studied. The PECVD oxide thickness was 0.5 µm and 0.3 µm on the front side and the back side of the wafers, respectively. Oxide is deposited on both sides of the wafer to minimize wafer bow during polishing and improve planarization. A soft polish was performed to remove about 30 nm of the oxide and to smooth the front oxide surface originally having a root mean square of the micro-roughness (RMS) of ~0.56 nm to a final ~0.18 nm. A modified RCA1 solution was used to clean the wafer surfaces followed by spin-drying.

Two wafers were loaded into the plasma system, both wafers are placed on the RF electrode and treated in plasma in RIE mode. For comparison, some wafers were treated in plasma mode in which the wafers were put on the grounded electrode. An oxygen plasma was used with a nominal flow rate of 16 scc/m. The RF power was 20-400 W (typically 80 W) at 13.56 MHz and the vacuum level was 100 mTorr. The oxide covered wafers were treated in plasma for times between 15 seconds to 5 minutes. The plasma treated silicon wafers were then dipped in an appropriate solution or rinse with de-ionized water followed by spin-drying and room temperature bonding in air. Some of the plasma treated wafers were also directly bonded in air without rinse or dipping.

The bonding energy was measured by inserting a wedge into the interface to measure the crack length according to the equation:

$$\gamma = \frac{3t_b^2 E_1 t_{w1}^3 E_2 t_{tw2}^3}{16L^4 (E_1 t_{w1}^3 + E_2 t_{w2}^3)}$$

E and two are the Young's modulus and thickness for wafers one and two and the is the thickness of a wedge inserted between the two wafers that results in a wafer separation of length L from the edge of the wafers.

The room temperature bonding energy as a function of storage time of bonded plasma treated oxide covered silicon wafers is shown in FIG. 6A. This figure shows measured room temperature bonding energy versus storage time for 4 different cases as shown. The results can be summarized as follows: (1) for dipped and bonded RIE plasma treated oxide wafers, the room temperature bonding energy increases with storage time and reaches a stable value after ~20 h in air or at low vacuum; (2) RIE mode results in higher bonding energies than plasma mode; (3) too short a plasma exposure time or too low a plasma power provides a small or negligible increase in bond energy; (4) NH₄OH dip after plasma treatment shows a much higher increase in bonding energy than water rinse; (5) direct bonding in air after plasma treatment without dipping or rinse shows an almost constant bonding energy with time. The bonding energy of the directly bonded wafer pairs immediately after room temperature bonding is slightly higher than the de-ionized water rinsed or NH₄OH dipped wafer pairs.

FIG. 6B shows room temperature bonding of Si and AlN wafers with PECVD oxide deposited layers. After about 100 h of storage time a bonding energy of over 2000 mJ/m² were observed.

Comparing different bonding materials, the bonding energy as a function of storage time of O_2 plasma treated thermally oxidized silicon wafer pairs is similar to wafers with PECVD oxide, although the values of the room temperature bonding energy are somewhat lower.

After \sim 24 h storage in air at room temperature, the bonding energy as high as \sim 1000 mJ/m² was reached in the RIE mode plasma treated and NH₄OH dipped PECVD oxide covered wafer pairs. Since the maximum bonding energy of a van der Waals bonded silicon oxide covered wafer pairs is about 200 mJ/m², a large portion of the bonding energy is attributed to the formation of covalent bonds at the bonding interface at room temperature according to the above equation.

EXAMPLES 2-3

The above process was applied to bond processed InP wafers (600 µm thick) to AlN wafers (380 µm thick), or to bond processed Si (380 µm thick) and InP 600 µm thick) wafers, as second and third examples. The processed InP device wafers are covered with PECVD oxide and planarized and smoothed by chemical-mechanical polishing CMP. A PECVD oxide layer is also deposited on the AlN wafers and is planarized and smoothed to improve

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the RMS surface roughness. The processed Si and processed InP wafers are deposited with PECVD oxide and planarized and smoothed using CMP. After VSE similar to the example 1 bonding at room temperature, the bonded wafers are left in ambient air at room temperature.

After 24 hours storage at room temperature, bonding energy of 1000 mJ/m² and 1100 mJ/m² were achieved for the InP/Si and InP/AlN bonded pairs, respectively. For processed Si (380 µm thick) /oxide covered AlN (280 µm thick) wafer pairs, the bonding energy at room temperature as high as 2500 mJ/m² has been achieved. These room temperature bonded plasma treated wafer pairs have sufficient bonding strength to sustain subsequent substrate lapping and etching and other typical semiconductor fabrication processes before or after substrate removal.

The InP substrate in the room temperature bonded InP/AlN pairs was lapped with 1900# Al₂O₃ powder from initial 600 μ m thick to ~50 μ m thick followed by etching in an HCl/H₃PO₄ solution to leave about a 2.0 μ m thick InP device layer on the AlN or Si wafer. The water and etching solution did not penetrate into the bonding interface.

Surfaces are sputter etched by energetic particles such as radicals, ions, photons and electrons in the plasma or RIE mode. For example, the O_2 plasma under conditions that bring about the desired VSE is sputter-etching about 2 Å/min of PECVD oxide as measured by a reflectance spectrometry. For thermal oxide the sputter etching rate is about 0.5 Å/min. The thickness of oxide before and after plasma treatment was measured by a reflectance spectrometry and averaged from 98 measured points on each wafer. The etching by O_2 plasma has not only cleaned the surface by oxidation and sputtering but also broken bonds of the oxide on the wafer surfaces.

However, the surface roughness of plasma treated oxide surfaces must not be degraded by the etching process. AFM measurements show that compared with the initial surface roughness, the RMS of the O_2 plasma treated oxide wafers was \sim 2 Å and did not change noticeably. On the other hand, if the etching is not sufficiently strong, the bonding energy enhancement effect is also small. Keeping other conditions unchanged when the O_2 plasma treatment was performed with plasma mode rather than RIE mode, the etching of oxide surfaces is negligible and the oxide thickness does not change. The final room temperature bonding energy is only 385 mJ/m² compared to 1000 mJ/m² of RIE treated wafers (see FIG. 6A).

Other gas plasma has shown a similar effect. CF_4/O_2 RIE was used to remove ~4 nm of PECVD oxide from the wafer surfaces prior to bonding. The bonding energy of room

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temperature bonded PECVD oxide covered silicon wafers was also enhanced significantly in this manner and exceeds 1000 mJ/m² after sufficient storage time (see also FIG. 6A).

An argon plasma has also been used for the VSE with a nominal flow rate of 16 scc/m. The RF power was typically 60 W at 13.56 MHz and the vacuum level was 100 mTorr. The oxide covered silicon wafers were treated in plasma in RIE mode for times between 30 seconds to 2 minutes. The plasma treated silicon wafers were then dipped in an NH₄OH solution followed by spin-drying and room temperature bonding in air. The bonding energy reached $\sim 800 \text{ mJ/m}^2$ at room temperature after only 8 h storage in air.

Obviously, numerous modifications and variations of the present invention are possible in light of the above teachings. It is therefore to be understood that within the scope of the appended claims, the invention may be practiced otherwise than as specifically described herein.

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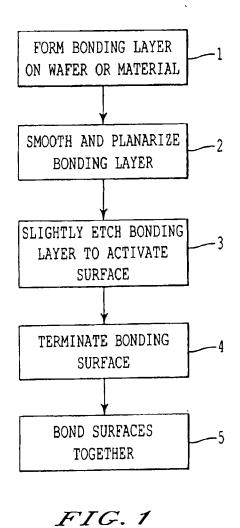
CLAIMS:

A bonding method, comprising:
 forming first and second bonding surfaces;
 etching said first and second bonding surfaces; and

bonding together at room temperature said first and second bonding surfaces after said etching step.

ABSTRACT OF THE INVENTION

A method for bonding at low or room temperature includes steps of surface cleaning and activation by cleaning or etching. The method may also include removing by-products of interface polymerization to prevent a reverse polymerization reaction to allow room temperature chemical bonding of materials such as silicon, silicon nitride and SiO₂. The surfaces to be bonded are polished to a high degree of smoothness and planarity. VSE may use reactive ion etching or wet etching to slightly etch the surfaces being bonded. The surface roughness and planarity are not degraded and may be enhanced by the VSE process. The etched surfaces may be rinsed in solutions such as ammonium hydroxide or ammonium fluoride to promote the formation of desired bonding species on the surfaces.



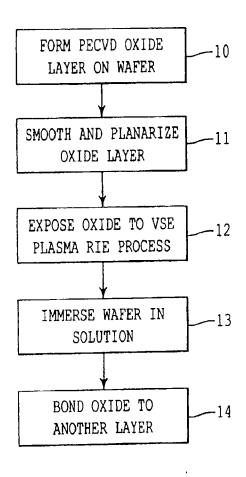
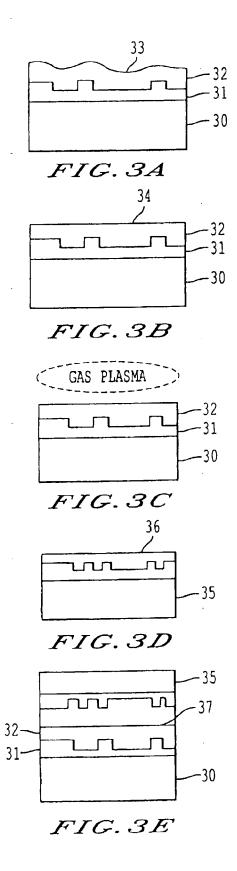
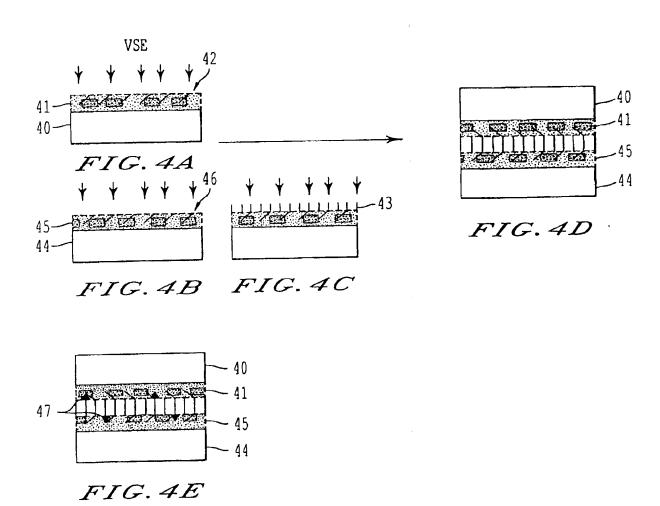
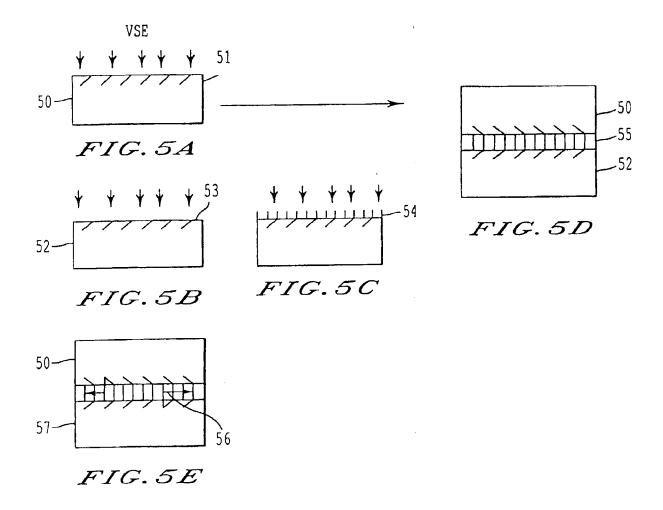
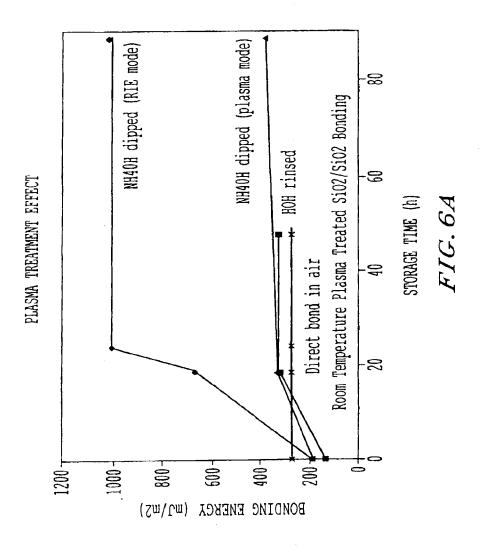


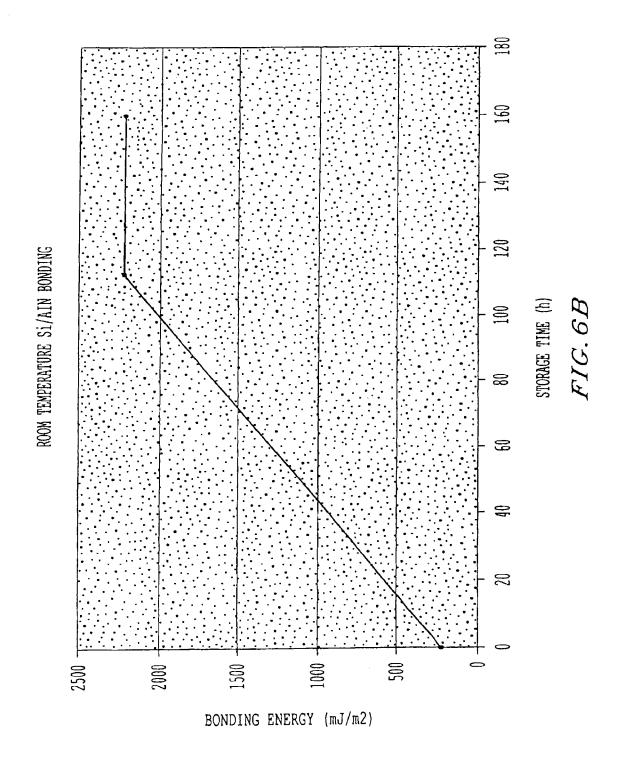
FIG.2

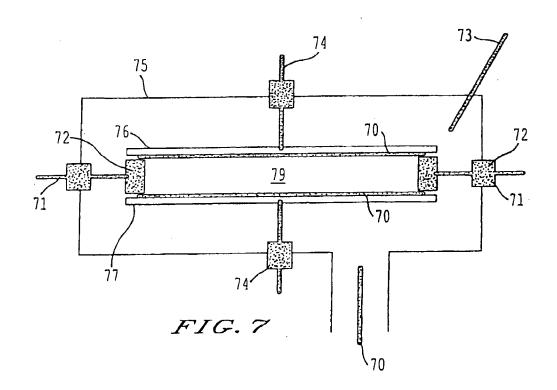


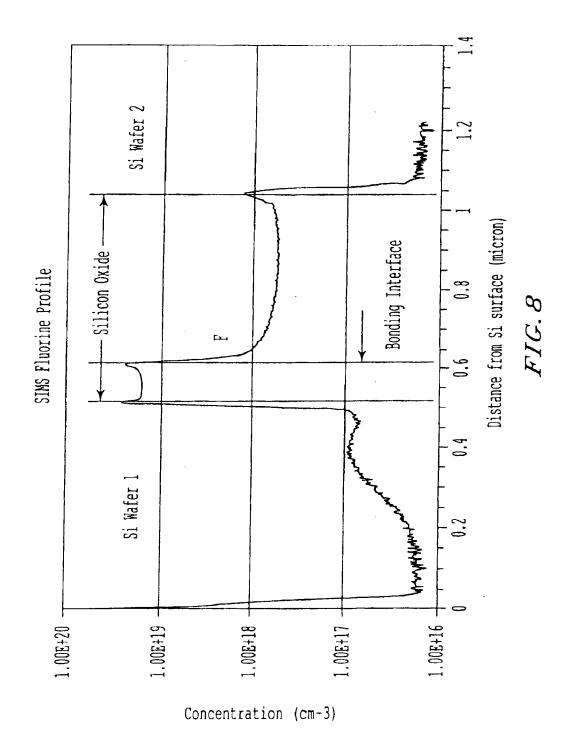












Doc Code: Oath

Document Description: Oath or declaration filed

SUBSTITUTE STATEMENT IN LIEU OF AN OATH OR DECLARATION FOR UTILITY OR DESIGN PATENT APPLICATION (35 U.S.C. 115(d) AND 37 CFR 1.64)

Title of Invention	METHOD FOR LOW TEMPERATURE BONDING AND BONDED STRUCTURE				
This statement is d	irected to:				
The attached a	application,				
OR					
United States	application or PCT international appl	ication number 14/19	7,070 filed on	03/04/2014 .	
LEGAL NAME o	f inventor to whom this substitute s	statement applies:			
(E.g., Given Name	(first and middle (if any)) and Family	y Name or Surname)			
Qin-Yi TONG	,				
Residence (except	for a deceased or legally incapacitate	d inventor):			
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				Country	
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I believe the above claimed invention	e-named inventor or joint inventor to in the application.	be the original inventor	or an original joint in	nventor of a	
The above-identifi	ied application was made or authorize	ed to be made by me.			
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Person to	whom the inventor is under an oblig	gation to assign,			
Person who otherwise shows a sufficient proprietary interest in the matter (petition under 37 CFR 1.46 is required), or					
	rentor.				

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	UBSITIUTE STATE	JVIEJN I	
Circumstances permitting execution of	this substitute statement:		
Inventor is deceased,			
Inventor is under legal incapacity	/ ,		
Inventor cannot be found or reac	hed after diligent effort, or		
Inventor has refused to execute t	he oath or declaration under 3	7 CFR 1.63.	
If there are joint inventors, please check	k the appropriate box below:		
An application data sheet under a entity has been or is currently sul	37 CFR 1.76 (PTO/AIA/14 or omitted.	equivalent) naming the	e entire inventive
OR			
An application data sheet under Substitute Statement Supplement and providing inventor informations.	tal Sheet (PTO/AIA/11 or equi	valent) naming the ent	en submitted. Thus, a ire inventive entity
PERSON EXECUTING THIS SUBS	STITUTE STATEMENT:		
Paul M. ENQUIST Name:			U //6/14 Date (Optional):
Signature: 12 M 5			
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Gary Purhan	North Carolina State	ファファ ナ 2 7513 Zip	United States Country
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DECLARATION (37 CFR 1.63) FOR UTILITY OR DESIGN APPLICATION USING AN APPLICATION DATA SHEET (37 CFR 1.76)

Title of Invention	METHOD FOR LOW TEMPERATURE BONDING AND BONDED STRUCTURE
As a below named i	nventor, I hereby declare that:
This declaration is o	directed to:
□т	he attached application, or
	nited States application or PCT international application number 4/197,070 Filed on MARCH 4, 2014
The above-identifie	d application was made or authorized to be made by me.
I believe that I am t application.	he original inventor or an original joint inventor of a claimed invention in the
	dge that any willful false statement made in this declaration is punishable 01 by fine or imprisonment of not more than five (5) years, or both.
LEGAL NAME OF	
	Gillman FOUNTAIN, Jr. Date (Optional): 4-16-2014
Signature:	us Hillran tourt fr.
	eet (PTO/SB/14 or equivalent), including naming the entire inventive entity, must accompany this form. form for each additional inventor.

DECLARATION (37 CFR 1.63) FOR UTILITY OR DESIGN APPLICATION USING AN APPLICATION DATA SHEET (37 CFR 1.76)

Title of Invention	METHOD FOR LOW TEMPERATURE BONDING AND BONDED STRUCTURE
As a below named i	nventor, I hereby declare that:
This declaration is o	directed to:
ПТ	he attached application, or
	Inited States application or PCT international application number 4/197,070 Filed on MARCH 4, 2014 .
The above-identifie	ed application was made or authorized to be made by me.
I believe that I am t application.	he original inventor or an original joint inventor of a claimed invention in the
I hereby acknowle under 18 U.S.C. 10	dge that any willful false statement made in this declaration is punishable 01 by fine or imprisonment of not more than five (5) years, or both.
LEGAL NAME OF	
Inventor: Paul M	1. ENQUIST Date (Optional): 4//6/14
Signature:	47571
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Application Da	ta Sheet 37 CF	R 1 76	Attorney	Docket N	umber	TSSRA.012C1		
Application Data Sheet 37 CFR 1.76		1.70	Application	on Numbe	er			
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Only complete this section when filing an application by reference under 35 U.S.C. 111(c) and 37 CFR 1.57(a). Do not complete this section if application papers including a specification and any drawings are being filed. Any domestic benefit or foreign priority information must be provided in the appropriate section(s) below (i.e., "Domestic Benefit/National Stage Information" and "Foreign Priority Information"). For the purposes of a filing date under 37 CFR 1.53(b), the description and any drawings of the present application are replaced by this reference to the previously filed application, subject to conditions and requirements of 37 CFR 1.57(a).										
Application number of the previously filed application filed appli						or Country				
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Application Da	ata Shaat 37 CED 1 76	Attorney Docket Number	TSSRA.012C1
Application Data Sheet 37 CFR 1.76		Application Number	
Title of Invention	METHOD FOR LOW TEMPE	RATURE BONDING AND BONE	DED STRUCTURE

Representative Information:

Representative information should be provided for all practitioners having a power of attorney in the application. Providing this information in the Application Data Sheet does not constitute a power of attorney in the application (see 37 CFR 1.32). Either enter Customer Number or complete the Representative Name section below. If both sections are completed the customer Number will be used for the Representative Information during processing.							
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Prior Applicati	on Status	Pending		v			Rer	nove
Application N	lumber	Conti	nuity Type		Prior Application Num	nber		or 371(c) Date YY-MM-DD)
		Continuation of	of	7	14/754111		2015-06-29	
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Application Number	Cont	inuity Type	Prior Applicati Number	on	Filing Date (YYYY-MM-DD)	Pa	tent Number	Issue Date (YYYY-MM-DD)
14/754111	Division o	f 🔻	14/197070		2014-03-04	90	82627	2015-07-14
Prior Applicati	on Status	Abandoned		7			Rer	nove
Application N	lumber	Conti	nuity Type		Prior Application Num	nber		or 371(c) Date YY-MM-DD)
14/197070		Division of		7	13/341273		2011-12-30	
Prior Applicati	on Status	Patented		7			Rer	nove
Application Number	Cont	inuity Type	Prior Applicati Number	on	Filing Date (YYYY-MM-DD)	Pa	tent Number	Issue Date (YYYY-MM-DD)
13/341273	Continuat	ion of	12/954740		2010-11-26	81	53505	2012-04-10
Prior Applicati	on Status	Patented		-			Rer	nove
Application Number	Cont	inuity Type	Prior Applicati Number	on	Filing Date (YYYY-MM-DD)	Pa	tent Number	Issue Date (YYYY-MM-DD)
12/954740	Continuat	ion of	12/720368		2010-03-09	78	71898	2011-01-18

Application Data Sheet 37 CFR 1.76					Attorney Docket Number		TSSRA.012C1			
					Application Number					
Title of Invention METHOD FOR LOW TEMPERATURE BONDING A					DING AND BONE	DED STR	UCTURE			
Prior Application	n Status	Patented			▼			Re	nove	
Application Number	··· I COMMUNICATION			Pri	or Application Number	Filing Da (YYYY-MM		Patent Number	Issue Date (YYYY-MM-DD)	
12/720368	Continuation	n of	-	11/98	30664	2007-10-31		7807549	2010-10-05	
Prior Application	n Status	Patented			•			Remove		
Application Number	Contin	uity Type		Pri	or Application Number	Filing Da (YYYY-MM		Patent Number	Issue Date (YYYY-MM-DD)	
11/980664	Continuation	n of	~	10/9	13441	2004-08-09		7387944	2008-06-17	
Prior Application Status Patented				•			Re	nove		
Application Continuity Type		Pri	or Application Number	Filing Date (YYYY-MM-DD)		Patent Number	Issue Date (YYYY-MM-DD)			
10/913441 Continuation of			05283	2000-02-16		6902987	2005-06-07			
Additional Domestic Benefit/National Stage Data may be generated within this form by selecting the Add button.										

Foreign Priority Information:

This section allows for the applicant to claim priority to a foreign application. Providing this information in the application data sheet constitutes the claim for priority as required by 35 U.S.C. 119(b) and 37 CFR 1.55. When priority is claimed to a foreign application that is eligible for retrieval under the priority document exchange program (PDX)ⁱ the information will be used by the Office to automatically attempt retrieval pursuant to 37 CFR 1.55(i)(1) and (2). Under the PDX program, applicant bears the ultimate responsibility for ensuring that a copy of the foreign application is received by the Office from the participating foreign intellectual property office, or a certified copy of the foreign priority application is filed, within the time period specified in 37 CFR 1.55(g)(1).

			Remove
Application Number	Country ⁱ	Filing Date (YYYY-MM-DD)	Access Code ⁱ (if applicable)
Additional Foreign Priority Add button.	Data may be generated wit	hin this form by selecting the	Add

Statement under 37 CFR 1.55 or 1.78 for AIA (First Inventor to File) Transition Applications

This application (1) claims priority to or the benefit of an application filed before March 16, 2013 and (2) also
contains, or contained at any time, a claim to a claimed invention that has an effective filing date on or after March
16, 2013.
NOTE: By providing this statement under 37 CFR 1.55 or 1.78, this application, with a filing date on or after March
16, 2013, will be examined under the first inventor to file provisions of the AIA.

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Application Da	ta Sheet 37 CFR 1.76	Attorney Docket Number	TSSRA.012C1
Application Da	ita Sileet Si Ci K 1.70	Application Number	
Title of Invention	METHOD FOR LOW TEMPER	RATURE BONDING AND BONE	DED STRUCTURE

Authorization or Opt-Out of Authorization to Permit Access:

When this Application Data Sheet is properly signed and filed with the application, applicant has provided written authority to permit a participating foreign intellectual property (IP) office access to the instant application-as-filed (see paragraph A in subsection 1 below) and the European Patent Office (EPO) access to any search results from the instant application (see paragraph B in subsection 1 below).

Should applicant choose not to provide an authorization identified in subsection 1 below, applicant <u>must opt-out</u> of the authorization by checking the corresponding box A or B or both in subsection 2 below.

NOTE: This section of the Application Data Sheet is **ONLY** reviewed and processed with the **INITIAL** filing of an application. After the initial filing of an application, an Application Data Sheet cannot be used to provide or rescind authorization for access by a foreign IP office(s). Instead, Form PTO/SB/39 or PTO/SB/69 must be used as appropriate.

- Authorization to Permit Access by a Foreign Intellectual Property Office(s)
- A. <u>Priority Document Exchange (PDX)</u> Unless box A in subsection 2 (opt-out of authorization) is checked, the undersigned hereby <u>grants the USPTO authority</u> to provide the European Patent Office (EPO), the Japan Patent Office (JPO), the Korean Intellectual Property Office (KIPO), the State Intellectual Property Office of the People's Republic of China (SIPO), the World Intellectual Property Organization (WIPO), and any other foreign intellectual property office participating with the USPTO in a bilateral or multilateral priority document exchange agreement in which a foreign application claiming priority to the instant patent application is filed, access to: (1) the instant patent application-as-filed and its related bibliographic data, (2) any foreign or domestic application to which priority or benefit is claimed by the instant application and its related bibliographic data, and (3) the date of filing of this Authorization. See 37 CFR 1.14(h) (1).
- B. <u>Search Results from U.S. Application to EPO</u> Unless box B in subsection 2 (opt-out of authorization) is checked, the undersigned hereby <u>grants the USPTO authority</u> to provide the EPO access to the bibliographic data and search results from the instant patent application when a European patent application claiming priority to the instant patent application is filed. See 37 CFR 1.14(h)(2).

The applicant is reminded that the EPO's Rule 141(1) EPC (European Patent Convention) requires applicants to submit a copy of search results from the instant application without delay in a European patent application that claims priority to the instant application.

2.	Opt-Out of Authorizations to Permit Access b	v a Foreign Intelle	ectual Property Office(s

A. Applicant <u>DOES NOT</u> authorize the USPTO to permit a participating foreign IP office access to the instant
application-as-filed. If this box is checked, the USPTO will not be providing a participating foreign IP office with
any documents and information identified in subsection 1A above.
D. A. II. ADDED NOT. III. III. HODEO A. A. III. III. EDO.

B. Applicant DOES NOT authorize the USPTO to transmit to the EPO any search results from the instant patent
application. If this box is checked, the USPTO will not be providing the EPO with search results from the instant
application.

NOTE: Once the application has published or is otherwise publicly available, the USPTO may provide access to the application in accordance with 37 CFR 1.14.

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Application Da	ita Sheet 37 CFR 1.76	Attorney Docket Number	TSSRA.012C1
Application Da	ita Sileet Si Ci K 1.70	Application Number	
Title of Invention	METHOD FOR LOW TEMPER	RATURE BONDING AND BONI	DED STRUCTURE

Applicant Information:

Providing assignment information in this section does not substitute for compliance with any requirement of part 3 of Title 37 of CFR to have an assignment recorded by the Office.							
Applicant 1			Remove				
If the applicant is the inventor (or the remaining joint inventor or inventors under 37 CFR 1.45), this section should not be completed. The information to be provided in this section is the name and address of the legal representative who is the applicant under 37 CFR 1.43; or the name and address of the assignee, person to whom the inventor is under an obligation to assign the invention, or person who otherwise shows sufficient proprietary interest in the matter who is the applicant under 37 CFR 1.46. If the applicant is an applicant under 37 CFR 1.46 (assignee, person to whom the inventor is obligated to assign, or person who otherwise shows sufficient proprietary interest) together with one or more joint inventors, then the joint inventor or inventors who are also the applicant should be identified in this section.							
Assignee	Legal Representative ur	nder 35 U.S.C. 117	Joint Inventor				
Person to whom the inventor is obli	gated to assign.	Person who show	vs sufficient proprietary interest				
If applicant is the legal representat	ive, indicate the authority to	file the patent application	on, the inventor is:				
			▼				
Name of the Deceased or Legally	Incapacitated Inventor:						
If the Applicant is an Organization	n check here.						
Organization Name	IX, INC.						
Mailing Address Information Fo	or Applicant:						
Address 1 800 F	erimeter Park Drive						
Address 2 Suite	В	_					
City	sville	State/Province	NC				
Country US		Postal Code	27560				
Phone Number	hone Number Fax Number						
Email Address							
Additional Applicant Data may be generated within this form by selecting the Add button.							

Assignee Information including Non-Applicant Assignee Information:

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Under the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it contains a valid OMB control number.

Application Data Sheet 37 CFR 1.76		Attorney Doo	ket Number	TSSRA.0)12C1				
Application	Dala Sii	ieel 37	CFK 1.76	Application N	lumber				
Title of Inventio	n METH	HOD FOR	R LOW TEMPER	RATURE BOND	ING AND BO	NDED STRU	CTURE		
Assignee 1	Assignee 1								
Complete this section if assignee information, including non-applicant assignee information, is desired to be included on the patent application publication. An assignee-applicant identified in the "Applicant Information" section will appear on the patent application bublication as an applicant. For an assignee-applicant, complete this section only if identification as an assignee is also desired on the patent application publication.									
If the Assignee	or Non-Ar	plicant	Assignee is an	Organization	check here.			Remove	
Prefix		Given N		Middle Nan	1	Family Na	me	Suffix	
	-								-
 Mailing Address	s Informat	tion Fo	r Assignee ind	Luding Non-	Applicant As	ssignee:			
Address 1									
Address 2									
City					State/Prov	ince			
Country					Postal Cod	ital Code			
Phone Number					Fax Numb	er			
Email Address					!	+			
Additional Assig selecting the Ad		n-Applic	cant Assignee	Data may be ç	generated wit	thin this forr	n by	Add]
Signature:								Remove]
NOTE: This Application Data Sheet must be signed in accordance with 37 CFR 1.33(b). However, if this Application Data Sheet is submitted with the INITIAL filing of the application and either box A or B is not checked in subsection 2 of the "Authorization or Opt-Out of Authorization to Permit Access" section, then this form must also be signed in accordance with 37 CFR 1.14(c). This Application Data Sheet must be signed by a patent practitioner if one or more of the applicants is a juristic entity (e.g., corporation or association). If the applicant is two or more joint inventors, this form must be signed by a patent practitioner, all joint inventors who are the applicant, or one or more joint inventor-applicants who have been given power of attorney (e.g., see USPTO Form PTO/AIA/81) on behalf of all joint inventor-applicants. See 37 CFR 1.4(d) for the manner of making signatures and certifications.									
Signature /Pa	aul Stellmai	n/				Date (Y	YYY-MM-DI	D) 2015	-12-02
First Name	Paul		Last Name	Stellman		Registra	ation Number	6828	7
Additional Signature may be generated within this form by selecting the Add button. Add									

U.S. Patent and Trademark Office; U.S. DEPARTMENT OF COMMERCE

Application Data Sheet 37 CFR 1.76		Attorney Docket Number	TSSRA.012C1	
		Application Number		
Title of Invention	METHOD FOR LOW TEMPERATURE BONDING AND BONDED STRUCTURE			

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This collection of information is required by 37 CFR 1.76. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to take 23 minutes to complete, including gathering, preparing, and submitting the completed application data sheet form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. **SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.**

Privacy Act Statement

The Privacy Act of 1974 (P.L. 93-579) requires that you be given certain information in connection with your submission of the attached form related to a patent application or patent. Accordingly, pursuant to the requirements of the Act, please be advised that: (1) the general authority for the collection of this information is 35 U.S.C. 2(b)(2); (2) furnishing of the information solicited is voluntary; and (3) the principal purpose for which the information is used by the U.S. Patent and Trademark Office is to process and/or examine your submission related to a patent application or patent. If you do not furnish the requested information, the U.S. Patent and Trademark Office may not be able to process and/or examine your submission, which may result in termination of proceedings or abandonment of the application or expiration of the patent.

The information provided by you in this form will be subject to the following routine uses:

- 1 The information on this form will be treated confidentially to the extent allowed under the Freedom of Information Act (5 U.S.C. 552) and the Privacy Act (5 U.S.C. 552a). Records from this system of records may be disclosed to the Department of Justice to determine whether the Freedom of Information Act requires disclosure of these records.
- 2. A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement negotiations.
- 3 A record in this system of records may be disclosed, as a routine use, to a Member of Congress submitting a request involving an individual, to whom the record pertains, when the individual has requested assistance from the Member with respect to the subject matter of the record.
- 4. A record in this system of records may be disclosed, as a routine use, to a contractor of the Agency having need for the information in order to perform a contract. Recipients of information shall be required to comply with the requirements of the Privacy Act of 1974, as amended, pursuant to 5 U.S.C. 552a(m).
- 5. A record related to an International Application filed under the Patent Cooperation Treaty in this system of records may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant to the Patent CooperationTreaty.
- 6. A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
- 7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or his/her designee, during an inspection of records conducted by GSA as part of that agency's responsibility to recommend improvements in records management practices and programs, under authority of 44 U.S.C. 2904 and 2906. Such disclosure shall be made in accordance with the GSA regulations governing inspection of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make determinations about individuals.
- 8. A record from this system of records may be disclosed, as a routine use, to the public after either publication of the application pursuant to 35 U.S.C. 122(b) or issuance of a patent pursuant to 35 U.S.C. 151. Further, a record may be disclosed, subject to the limitations of 37 CFR 1.14, as a routine use, to the public if the record was filed in an application which became abandoned or in which the proceedings were terminated and which application is referenced by either a published application, an application open to public inspections or an issued patent.
- A record from this system of records may be disclosed, as a routine use, to a Federal, State, or local law enforcement agency, if the USPTO becomes aware of a violation or potential violation of law or regulation.

Electronic Patent Application Fee Transmittal						
Application Number:						
Filing Date:						
Title of Invention:	ME	THOD FOR LOW TE	MPERATURE BOI	NDING AND BONI	DED STRUCTURE	
First Named Inventor/Applicant Name:	Qin	Qin-Yi Tong				
Filer: Paul Steven Stellman/Jonathan Pan						
Attorney Docket Number: TSSRA.012C1						
Filed as Large Entity	'					
Filing Fees for Utility under 35 USC 111(a)						
Description		Fee Code	Quantity	Amount	Sub-Total ir USD(\$)	
Basic Filing:			1			
Utility application filing		1011	1	280	280	
Utility Search Fee		1111	1	600	600	
Utility Examination Fee		1311	1	720	720	
Pages:	<u>'</u>		<u> </u>			
Claims:						
Miscellaneous-Filing:						
Petition:						
Patent-Appeals-and-Interference: age 45 of 331				SAMSLIN	G EXHIBIT 1	

Description	Fee Code	Quantity	Amount	Sub-Total in USD(\$)
Post-Allowance-and-Post-Issuance:				
Extension-of-Time:				
Miscellaneous:				
	Tot	al in USD	(\$)	1600

Electronic Acknowledgement Receipt				
EFS ID:	24246561			
Application Number:	14957501			
International Application Number:				
Confirmation Number:	1033			
Title of Invention:	METHOD FOR LOW TEMPERATURE BONDING AND BONDED STRUCTURE			
First Named Inventor/Applicant Name:	Qin-Yi Tong			
Customer Number:	135980			
Filer:	Paul Steven Stellman/Tony Do			
Filer Authorized By:	Paul Steven Stellman			
Attorney Docket Number:	TSSRA.012C1			
Receipt Date:	02-DEC-2015			
Filing Date:				
Time Stamp:	20:20:34			
Application Type:	Utility under 35 USC 111(a)			

Payment information:

Submitted with Payment	yes
Payment Type	Credit Card
Payment was successfully received in RAM	\$1600
RAM confirmation Number	20724
Deposit Account	
Authorized User	

The Director of the USPTO is hereby authorized to charge indicated fees and credit any overpayment as follows:

File Lietie e									
Document Number	Document Description	File Name	File Size(Bytes)/ Message Digest	Multi Part /.zip	Pages (if appl.)				
Number			153218	rait/.zip	(ii appi.)				
1		2015-12-02_Specification_TSSR A-012C1.pdf		yes	22				
			4914a5a4834012191f2fc21c18937e08afbb 3406						
Multipart Description/PDF files in .zip description									
	Document De	Start	E	nd					
	Specificat	ion	1	2	20				
	Claims	21	2	<u>?</u> 1					
	Abstrac	22 22							
Warnings:									
Information:									
2	Drawings-only black and white line	2015-12-02_Drawings_TSSRA-0	110488	no	9				
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Information:									
3	Oath or Declaration filed	2015-12-02_Declaration_TSSRA	142370	no	4				
3	Oath of Declaration flied	-012C1.pdf	cea2c9f3edf38cff13712c00f26b1f18dcb14 1dc	no	4				
Warnings:									
Information:									
4	Application Data Sheet	2015-12-02_ADS_TSSRA-012C1	1823736		9				
7	Application Data Sheet	.pdf	h7289574b97f38928edf98b24674de618eb d1170		,				
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5	Fee Worksheet (SB06)	fee-info.pdf	35336	no	2				
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Warnings:									
Information:									
		Total Files Size (in bytes):	22	65148					

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New Applications Under 35 U.S.C. 111

If a new application is being filed and the application includes the necessary components for a filing date (see 37 CFR 1.53(b)-(d) and MPEP 506), a Filing Receipt (37 CFR 1.54) will be issued in due course and the date shown on this Acknowledgement Receipt will establish the filing date of the application.

National Stage of an International Application under 35 U.S.C. 371

If a timely submission to enter the national stage of an international application is compliant with the conditions of 35 U.S.C. 371 and other applicable requirements a Form PCT/DO/EO/903 indicating acceptance of the application as a national stage submission under 35 U.S.C. 371 will be issued in addition to the Filing Receipt, in due course.

New International Application Filed with the USPTO as a Receiving Office

If a new international application is being filed and the international application includes the necessary components for an international filing date (see PCT Article 11 and MPEP 1810), a Notification of the International Application Number and of the International Filing Date (Form PCT/RO/105) will be issued in due course, subject to prescriptions concerning national security, and the date shown on this Acknowledgement Receipt will establish the international filing date of the application.

Index the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it displays a valid OMR control number

PA	PATENT APPLICATION FEE DETERMINATION RECORD Substitute for Form PTO-875 Application or Docket Number 14/957,501 Filing Date 12/02/2015 To be Mailed							
							ENTITY:	LARGE SMALL MICRO
				APPLICA	ATION AS FILE	D – PART	· I	
			(Column ·	1)	(Column 2)			
ᄂ	FOR	N	IUMBER FIL	_ED	NUMBER EXTRA	_	RATE (\$)	FEE (\$)
Ш	BASIC FEE (37 CFR 1.16(a), (b), (or (c))	N/A		N/A		N/A	
	SEARCH FEE (37 CFR 1.16(k), (i), c	or (m))	N/A		N/A		N/A	
	EXAMINATION FE (37 CFR 1.16(o), (p),		N/A		N/A		N/A	
	TAL CLAIMS CFR 1.16(i))		mir	nus 20 = *			X \$ =	
	EPENDENT CLAIM CFR 1.16(h))	S	m	inus 3 = *			X \$ =	
	If the specification and drawings exceed 100 sheets of paper, the application size fee due is \$310 (\$155 for small entity) for each additional 50 sheets or fraction thereof. See 35 U.S.C. 41(a)(1)(G) and 37 CFR 1.16(s).							
	MULTIPLE DEPEN	IDENT CLAIM PF	RESENT (3	7 CFR 1.16(j))				
* If t	he difference in colu	ımn 1 is less thar	zero, ente	r "0" in column 2.			TOTAL	
		(Column 1)		(Column 2)	(Column 3)	ED – PAI	RT II	
LN:	12/02/2015	CLAIMS REMAINING AFTER AMENDMENT		HIGHEST NUMBER PREVIOUSLY PAID FOR	PRESENT EXTE	RA	RATE (\$)	ADDITIONAL FEE (\$)
AMENDMENT	Total (37 CFR 1.16(i))	* 33	Minus	** 33	= 0		x \$80 =	0
N N	Independent (37 CFR 1.16(h))	* 4	Minus	***4	= 0		x \$420 =	0
AM	Application Si	ze Fee (37 CFR	1.16(s))					
	FIRST PRESEN	ITATION OF MULTI	PLE DEPEN	DENT CLAIM (37 CFF	R 1.16(j))			
							TOTAL ADD'L F	EE 0
		(Column 1)		(Column 2)	(Column 3)			
Τ		CLAIMS REMAINING AFTER AMENDMENT		HIGHEST NUMBER PREVIOUSLY PAID FOR	PRESENT EXTE	RA	RATE (\$)	ADDITIONAL FEE (\$)
ENT	Total (37 CFR 1.16(i))	*	Minus	**	=		X \$ =	
ENDM	Independent (37 CFR 1.16(h))	*	Minus	***	=		X \$ =	
	Application Si	ze Fee (37 CFR	1.16(s))					
AM	FIRST PRESEN	ITATION OF MULTI	PLE DEPEN	DENT CLAIM (37 CFF	R 1.16(j))			
Г	-						TOTAL ADD'L F	EE
** If ***	the entry in column the "Highest Numbe f the "Highest Numb	er Previously Paid per Previously Pai	l For" IN Th d For" IN T	HIS SPACE is less HIS SPACE is less	than 20, enter "20".		LIE /KIMBERLY	

This collection of information is required by 37 CFR 1.16. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to take 12 minutes to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS

ADDRESS. **SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.**

If you need assistance in completing the form, call 1-800-PTO-9199 and select option 2.

TSSRA.012C1 PATENT

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Inventor : Tong et al.

App. No. : 14,957,501

Filed: December 2, 2015

For : METHOD FOR LOW

TEMPERATURE BONDING AND

BONDED STRUCTURE

Examiner : Unknown

Art Unit : Unknown

Conf. No. : 1033

PRELIMINARY AMENDMENT

Mail Stop Amendment

Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450

Dear Sir:

Prior to examination on the merits, Applicant submits the following amendments.

Amendments to the Claims are reflected in the listing of claims which begins on page 2 of this paper.

Remarks/Arguments begin on page 6 of this paper.

Filing Date: December 2, 2015

AMENDMENTS TO THE CLAIMS

Please cancel Claim 1 without prejudice or disclaimer. Please add new Claims 2-34.

1. (Canceled)

2. (New) A bonding method, comprising:

forming a planarized insulating material on a semiconductor wafer;

etching the planarized insulating material in a reaction space;

terminating the planarized insulating material with a nitrogen-containing species by at least one of:

a nitrogen-containing plasma during the etching, and

a nitrogen-containing solution after the etching;

bringing a first material into direct contact with the terminated and planarized insulating material outside the reaction space; and

forming a chemical bond between the terminated and planarized insulating material and the first material.

- 3. (New) The method of Claim 2, wherein the etching the planarized insulating material is conducted in a vacuum chamber, the reaction space comprising the vacuum chamber.
- 4. (New) The method of Claim 2, wherein the etching the planarized insulating material is conducted in a plasma chamber, the reaction space comprising the plasma chamber.
- 5. (New) The method of Claim 2, wherein the etching and the terminating comprise exposing the planarized insulating material to the plasma etching process in a plasma chamber, the reaction space comprising the plasma chamber.
- 6. (New) The method of Claim 5, wherein the exposing the planarized insulating material to the plasma etching process comprises exposing the planarized insulating material to a reactive ion etching process.
- 7. (New) The method of Claim 5, wherein the exposing the planarized insulating material to the plasma etching process in the plasma chamber comprises using nitrogen gas.
- 8. (New) The method of Claim 2, wherein the terminating comprises exposing the planarized insulating material to a nitrogen-containing solution after the etching.
- 9. (New) The method of Claim 8, wherein the nitrogen-containing solution comprises an ammonia-based solution.

Filing Date: December 2, 2015

10. (New) The method of Claim 2, wherein the first material comprises a second planarized insulating material on a second semiconductor wafer.

- 11. (New) The method of Claim 10, wherein the first material comprises silicon oxide.
- 12. (New) The method of Claim 2, wherein the bringing into direct contact comprises bringing into direct contact the terminated planarized insulating material and a silicon material having a native oxide, the first material comprising the silicon material.
- 13. (New) The method of Claim 2, further comprising, after the terminating, rinsing the insulating material.
- 14. (New) The method of Claim 13, wherein the rinsing comprises immersing the insulating material in deionized water.
- 15. (New) The method of Claim 2, wherein the bringing into direct contact is performed at room temperature.
- 16. (New) The method of Claim 2, wherein the bringing into direct contact is conducted in air.
- 17. (New) The method of Claim 2, wherein the forming the planarized insulating material comprises forming silicon oxide on the semiconductor wafer.
- 18. (New) The method of Claim 2, further comprising forming a bond between the planarized insulating material and the first material with a strength of at least 500 mJ/m² without annealing at more than about 200° C.
- 19. (New) The method of Claim 2, further comprising forming a bond between the planarized insulating material and the first material with a strength of at least 2000 mJ/m^2 without annealing at more than about 200° C.
- 20. (New) The method of Claim 2, further comprising annealing the planarized insulating material and the first material at a temperature of no more than about 200° C after the bringing into direct contact.
- 21. (New) The method of Claim 2, wherein the planarized insulating material has a surface roughness between about 0.5 and 1.5 nm after the etching and before the bringing into direct contact.
 - 22. (New) A bonded structure comprising:

Filing Date: December 2, 2015

a first semiconductor material comprising a planarized insulating material having a first bonding surface; and

a second material having a second bonding surface,

wherein the first bonding surface is an etched surface terminated with a nitrogencontaining species, and

wherein the first and second bonding surfaces are in direct contact with each other and bonded together with a chemical bond without any intervening adhesive.

- 23. (New) The structure of Claim 22, further comprising at least one integrated circuit in the first semiconductor material.
 - 24. (New) The structure of Claim 22, wherein the second material comprises silicon.
- 25. (New) The structure of Claim 22, wherein the chemical bond has a strength of at least 500 mJ/m².
- 26. (New) The structure of Claim 22, wherein the chemical bond has a strength of at least 2000 mJ/m².
- 27. (New) The structure of Claim 22, wherein the planarized insulating material comprises silicon oxide.
- 28. (New) The structure of Claim 22, wherein the second bonding surface comprises an etched surface.
- 29. (New) The structure of Claim 28, wherein the etched surface of the second bonding surface is terminated with a nitrogen-containing species.
- 30. (New) The structure of Claim 22, wherein the first bonding surface is terminated with a nitrogen-containing species by at least one of a nitrogen-containing plasma during etching and a nitrogen-containing solution after etching.
 - 31. (New) The structure of Claim 22, wherein the chemical bond is a covalent bond.
 - 32. (New) A bonding method, comprising:

forming a planarized insulating material on a semiconductor wafer;

etching the planarized insulating material;

terminating the planarized insulating material with a nitrogen-containing species by at least one of:

a nitrogen-containing plasma during the etching, and

a nitrogen-containing solution after the etching;

Filing Date: December 2, 2015

bringing a first material into direct contact with the terminated and planarized insulating material; and

forming a chemical bond between the terminated and planarized insulating material and the first material.

33. (New) A bonding method, comprising:

forming a planarized insulating material on a processed semiconductor wafer;

exposing the planarized insulating material to a plasma process in a plasma chamber using a nitrogen-containing gas;

terminating the planarized insulating material with a species;

bringing into direct contact the terminated and planarized insulating material and a first material outside of said plasma chamber; and

forming a chemical bond between the terminated and planarized insulating material and the first material.

34. (New) The bonding method of Claim 33, wherein the species comprises nitrogen.

Filing Date: December 2, 2015

REMARKS

Prior to examination on the merits, Applicant has canceled Claim 1 without prejudice or

disclaimer. Claims 2-34 are new claims. Support for new Claims 2-34 may be found throughout

the Application as filed. No new matter is added. Applicant respectfully requests entry and

examination of the claims submitted herewith.

No Disclaimers or Disavowals

Although the present communication may include alterations to the application or claims,

Applicant is not conceding in this application that previously pending claims are not patentable.

Rather, any alterations or characterizations are being made to facilitate expeditious prosecution

of this application. Applicant reserves the right to pursue at a later date any previously pending

or other broader or narrower claims that capture any subject matter supported by the present

disclosure, including subject matter found to be specifically disclaimed herein or by any prior

prosecution. Accordingly, reviewers of this or any parent, child or related prosecution history

shall not reasonably infer that Applicant has made any disclaimers or disavowals of any subject

matter supported by the present application.

Please charge any additional fees, including any fees for additional extension of time, or

credit overpayment to Deposit Account No. 11-1410.

Respectfully submitted,

KNOBBE, MARTENS, OLSON & BEAR, LLP

Dated: December 3, 2015

By: /Paul Stellman/

Paul Stellman

Registration No. 68,287

Attorney of Record

Customer No. 135,980

(415) 954-4114

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-6-

Electronic Patent Application Fee Transmittal					
Application Number:	149	57501			
Filing Date:					
Title of Invention:	METHOD FOR LOW TEMPERATURE BONDING AND BONDED STRUCTURE				
First Named Inventor/Applicant Name:	Qin-Yi Tong				
Filer:	Paul Steven Stellman/Jonathan Pan				
Attorney Docket Number:	TSS	RA.012C1			
Filed as Large Entity					
Filing Fees for Utility under 35 USC 111(a)					
Description		Fee Code	Quantity	Amount	Sub-Total in USD(\$)
Basic Filing:					
Pages:					
Claims:					
Claims in Excess of 20		1202	13	80	1040
Independent claims in excess of 3		1201	1	420	420
Miscellaneous-Filing:					
Petition:					
Patent-Appeals-and-Interference:					

Description	Fee Code	Quantity	Amount	Sub-Total in USD(\$)
Post-Allowance-and-Post-Issuance:				
Extension-of-Time:				
Miscellaneous:				
	Tot	al in USD	(\$)	1460

Electronic Acknowledgement Receipt				
EFS ID:	24253209			
Application Number:	14957501			
International Application Number:				
Confirmation Number:	1033			
Title of Invention:	METHOD FOR LOW TEMPERATURE BONDING AND BONDED STRUCTURE			
First Named Inventor/Applicant Name:	Qin-Yi Tong			
Customer Number:	135980			
Filer:	Paul Steven Stellman/Heather OBrien			
Filer Authorized By:	Paul Steven Stellman			
Attorney Docket Number:	TSSRA.012C1			
Receipt Date:	03-DEC-2015			
Filing Date:				
Time Stamp:	15:53:18			
Application Type:	Utility under 35 USC 111(a)			

Payment information:

Submitted with Payment	yes
Payment Type	Credit Card
Payment was successfully received in RAM	\$1460
RAM confirmation Number	2422
Deposit Account	111410
Authorized User	KNOBBE MARTENS OLSON AND BEAR

The Director of the USPTO is hereby authorized to charge indicated fees and credit any overpayment as follows:

Charge any Additional Fees required under 37 C.F.R. Section 1.16 (National application filing, search, and examination fees)

File Listing: **Document** File Size(Bytes)/ Multi **Pages Document Description File Name** Number Message Digest Part /.zip (if appl.) 107322 2015-12-03_PreliminaryAmend 1 6 yes ment_TSSRA-012C1.pdf a2c9f923b2adf98105eff4d03e2417adf6d5 Multipart Description/PDF files in .zip description **Document Description** Start End Preliminary Amendment 1 1 5 Claims 2 Applicant Arguments/Remarks Made in an Amendment 6 6 Warnings: Information: 32396 2 Fee Worksheet (SB06) fee-info.pdf nο 2 637f62ff6fba759d3c2d23967261295925de 4465

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National Stage of an International Application under 35 U.S.C. 371

If a timely submission to enter the national stage of an international application is compliant with the conditions of 35 U.S.C. 371 and other applicable requirements a Form PCT/DO/EO/903 indicating acceptance of the application as a national stage submission under 35 U.S.C. 371 will be issued in addition to the Filing Receipt, in due course.

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Warnings:
Information:

139718

Doc Code: PA.,

Document Description: Power of Attorney

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POWER OF ATTORNEY BY APPLICANT

	y revoke ali pro ces below.	evious powers of attorney giver	in the applicat	ion identified in	<u>either</u> the a	ettached trai	nsmittal letter or
		Application Number		Filing Date	***************************************		
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lam the	Applicant (if the	Applicant is a juristic entity, list the	Applicant name	in the box):			W.
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		son to Whom the Inventor is Unde nerwise Shows Sufficient Proprieta					
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The L	indersigned (who	se julje is Jupplijid below) is authorize	ed to act on behalf	of the applicant (e	.g., where th	ne applicant is	a juristic entity).
Signa	***************************************	44 476		Date (Opti	onal)		
Name		Paul Davis V V		***************************************		***************************************	***************************************
Title		Senior Vice President, General	***************************************	***************************************	******	*****	·····
NOTE and c	Signature - Thi ertifications. If mo	s form must be signed by the applica- ire than one applicant, use multiple fo	ant in accordance or orms.	with 37 CFR 1.33.	See 37 CFR	t 1.4 for signat	ure requirements
Total	of	forms are submitted.					

This collection of information is required by 37 CFR 1.131, 1.32, and 1.33. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.11 and 1.14. This collection is estimated to take 3 minutes to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will very depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Petent and Trademark Office, U.S. Department of Commerce, P.O. Sox 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Sox 1450, Alexandria, VA 22313-1450.

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Electronic Acl	knowledgement Receipt
EFS ID:	24256974
Application Number:	14957501
International Application Number:	
Confirmation Number:	1033
Title of Invention:	METHOD FOR LOW TEMPERATURE BONDING AND BONDED STRUCTURE
First Named Inventor/Applicant Name:	Qin-Yi Tong
Customer Number:	135980
Filer:	Paul Steven Stellman/Gustavo Lopez
Filer Authorized By:	Paul Steven Stellman
Attorney Docket Number:	TSSRA.012C1
Receipt Date:	03-DEC-2015
Filing Date:	
Time Stamp:	17:06:12
Application Type:	Utility under 35 USC 111(a)

Payment information:

File Listing:

Document Number	Document Description	File Name	File Size(Bytes)/ Message Digest	Multi Part /.zip	Pages (if appl.)
1	Miscellaneous Incoming Letter	2015-12-03_Rescission_TSSRA- 012C1.pdf	64176 	no	1
Warnings:					

2	2015-12-03_POA_TSSRA-01: .pdf	2C1 246826 0d2e2770c5b94734e48a1aaa16ce4c0d0e6 df777	yes 2
	Multipart Description/PDF files	in .zip description	•
	Document Description	Start	End
	Transmittal Letter	1	1
	Power of Attorney	2	2

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Total Files Size (in bytes): 311002

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National Stage of an International Application under 35 U.S.C. 371

If a timely submission to enter the national stage of an international application is compliant with the conditions of 35 U.S.C. 371 and other applicable requirements a Form PCT/DO/EO/903 indicating acceptance of the application as a national stage submission under 35 U.S.C. 371 will be issued in addition to the Filing Receipt, in due course.

New International Application Filed with the USPTO as a Receiving Office

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Docket No.: TSSRA.012C1 December 3, 2015 App. No.: 14/957,501 Page 1 of 1

Please Direct All Correspondence to Customer Number 135,980

RESCISSION OF ANY PRIOR DISCLAIMERS AND REQUEST TO REVISIT ART

Inventor : Tong et al.

App. No : 14/957,501

Filed: December 2, 2015

For : METHOD FOR LOW

TEMPERATURE BONDING AND

BONDED STRUCTURE

Examiner : Unknown

Art Unit : Unknown

Conf # : 1033

Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450

Dear Sir:

The claims of the present application are different and possibly broader in scope than the claims pursued in the parent application(s). To the extent any prior amendments or characterizations of the scope of any claim or referenced art could be construed as a disclaimer of any subject matter supported by the present disclosure, Applicant hereby rescinds and retracts such disclaimer. Accordingly, the references previously considered in the parent application(s) may need to be re-visited.

Please charge any additional fees, including any fees for additional extension of time, or credit overpayment to Deposit Account No. 11-1410.

Knobbe, Martens, Olson & Bear, LLP Respectfully submitted,

Dated: December 3, 2015 /Paul Stellman/

Paul Stellman Registration No. 68,287 Attorney of Record Customer No. 135,980 (415) 954-4114

22195876//120315

PTO/AIA/82A (07-13) Approved for use through 11/30/2014. OMB 0651-0051

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Filing Date		December 2, 2015					
First Named Inve	Tong et al.						
Title		METHOD FOR LOW TEMPERATURE E STRUCTURE	BONDING AN	D BONDED			
Art Unit		Unknown					
Examiner Name		Unknown					
Attorney Docket Number		TSSRA.012C1					
SIGNATU	JRE of A	oplicant or Patent Practitioner					
Signature	/Paul St	ellman/	Date (Optional)	December 3, 2015			
Name Paul Ste		ellman	Registration Number	68,287			
Title (if Applicant is a juristic entity)							
Applicant Name (if Applicant is a juristic entity) NOTE: This form must be signed in accordance with 37 CFR 1.33. See 37 CFR 1.4(d) for signature requirements and certifications. If							
more than one applica							
*Total of 1 forms are submitted.							

This collection of information is required by 37 CFR 1.131, 1.32, and 1.33. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.11 and 1.14. This collection is estimated to take 3 minutes to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.

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APPLICATION NUMBER FILING OR 371(C) DATE FIRST NAMED APPLICANT ATTY. DOCKET NO./TITLE

14/957,501 12/02/2015 Qin-Yi Tong

TSSRA.012C1
CONFIRMATION NO. 1033

135980 Knobbe, Martens, Olson & Bear, LLP 2040 main Street Fourteenth Floor Irvine, CA 92614



Date Mailed: 12/15/2015

NOTICE OF ACCEPTANCE OF AUTHORIZATION TO PERMIT ACCESS TO APPLICATION VIA PRIORITY DOCUMENT EXCHANGE

This is in response to the applicant's authorization to permit access to the application-as-filed by participating offices under 37 CFR 1.14(h)(1) submitted on 12/02/2015.

The authorization to permit access to the application under 37 CFR 1.14(h)(1) is accepted.

Questions about the contents of this notice and the requirements it sets forth should be directed to the Office of Data Management, Application Assistance Unit, at (571) 272-4000 or (571) 272-4200 or 1-888-786-0101.

/fhadera/	



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APPLICATION NUMBER FILING OR 371(C) DATE FIRST NAMED APPLICANT ATTY. DOCKET NO./TITLE 14/957.501 12/02/2015 Qin-Yi Tong TSSRA.012C1

CONFIRMATION NO. 1033

135980 Knobbe, Martens, Olson & Bear, LLP 2040 main Street Fourteenth Floor Irvine, CA 92614



Date Mailed: 12/15/2015

NOTICE OF ACCEPTANCE OF AUTHORIZATION TO PERMIT ACCESS TO SEARCH RESULTS

This is in response to the applicant's authorization to permit access to the search results from the instant application under 37 CFR 1.14(h)(2) submitted on 12/02/2015.

The authorization to permit access to the search results under 37 CFR 1.14(h)(2) is accepted.

Questions about the contents of this notice and the requirements it sets forth should be directed to the Office of Data Management, Application Assistance Unit, at (571) 272-4000 or (571) 272-4200 or 1-888-786-0101.

/fhadera/

	PATEN	NT APPLI		ON FEE DE		ION RECORI	D	Applica 14/95	tion or Docket Num 7,501	nber
	APPLIC	CATION A		D - PART I	umn 2)	SMALL	ENTITY	OR	OTHER SMALL	
FOR NUMBER FILED NUMBER EXTRA				RATE(\$)	FEE(\$)		RATE(\$)	FEE(\$)		
	SIC FEE FR 1.16(a), (b), or (c))	N	/A	N	I/A	N/A		1	N/A	280
SEA	ARCH FEE FR 1.16(k), (i), or (m))	N	/A	N	I/A	N/A		1	N/A	600
ΞXΑ	MINATION FEE FR 1.16(o), (p), or (q))	N	/A	N	I/A	N/A		1	N/A	720
ОТ	AL CLAIMS FR 1.16(i))	33	minus	20= *	13			OR	x 80 =	1040
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FIRST NAMED APPLICANT ATTY. DOCKET NO./TITLE APPLICATION NUMBER FILING OR 371(C) DATE 14/957,501 12/02/2015 Qin-Yi Tong TSSRA.012C1

> **CONFIRMATION NO. 1033 INFORMAL NOTICE**

135980 Knobbe, Martens, Olson & Bear, LLP 2040 main Street Fourteenth Floor Irvine, CA 92614



Date Mailed: 12/15/2015

INFORMATIONAL NOTICE TO APPLICANT

Applicant is notified that the above-identified application contains the deficiencies noted below. No period for reply is set forth in this notice for correction of these deficiencies. However, if a deficiency relates to the inventor's oath or declaration, the applicant must file an oath or declaration in compliance with 37 CFR 1.63, or a substitute statement in compliance with 37 CFR 1.64, executed by or with respect to each actual inventor no later than the expiration of the time period set in the "Notice of Allowability" to avoid abandonment. See 37 CFR 1.53(f).

The item(s) indicated below are also required and should be submitted with any reply to this notice to avoid further processing delays.

- A properly executed inventor's oath or declaration has not been received for the following inventor(s): Qin-Yi Tong
- The substitute statement is signed by a person who is not identified as the applicant under 37 CFR 1.46 in an application data sheet (ADS). Only an applicant under 37 CFR 1.43, 1.45 or 1.46 may execute a substitute statement. See 37 CFR 1.64(a). An application data sheet in compliance with 37 CFR 1.76 identifying the person (i.e., assignee, obligated assignee, or person who otherwise shows sufficient proprietary interest) executing the substitute statement as the applicant, or an inventor's oath or declaration in compliance with 37 CFR 1.63 or 1.64 executed by or with respect to the inventor for whom the substitute statement was provided, is required.

Qin-Yi Tong

Questions about the contents of this notice and the requirements it sets forth should be directed to the Office of Data Management, Application Assistance Unit, at (571) 272-4000 or (571) 272-4200 or 1-888-786-0101.

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APPLICATION	FILING or	GRP ART				
NUMBER	371(c) DATE	UNIT	FIL FEE REC'D	ATTY.DOCKET.NO	TOT CLAIMS	IND CLAIMS
14/957,501	12/02/2015		3200	TSSRA.012C1	33	4

CONFIRMATION NO. 1033 FILING RECEIPT

135980 Knobbe, Martens, Olson & Bear, LLP 2040 main Street Fourteenth Floor Irvine, CA 92614



Date Mailed: 12/15/2015

Receipt is acknowledged of this non-provisional patent application. The application will be taken up for examination in due course. Applicant will be notified as to the results of the examination. Any correspondence concerning the application must include the following identification information: the U.S. APPLICATION NUMBER, FILING DATE, NAME OF APPLICANT, and TITLE OF INVENTION. Fees transmitted by check or draft are subject to collection. Please verify the accuracy of the data presented on this receipt. If an error is noted on this Filing Receipt, please submit a written request for a Filing Receipt Correction. Please provide a copy of this Filing Receipt with the changes noted thereon. If you received a "Notice to File Missing Parts" for this application, please submit any corrections to this Filing Receipt with your reply to the Notice. When the USPTO processes the reply to the Notice, the USPTO will generate another Filing Receipt incorporating the requested corrections

Inventor(s)

Qin-Yi Tong, Durham, NC;

Gaius Gillman Fountain JR., Youngsville, NC;

Paul M. Enquist, Cary, NC;

Applicant(s)

ZIPTRONIX, INC., Morrisville, NC;

Power of Attorney: The patent practitioners associated with Customer Number 135980

Domestic Priority data as claimed by applicant

This application is a CON of 14/754,111 06/29/2015 which is a DIV of 14/197,070 03/04/2014 PAT 9082627

which is a DIV of 13/341,273 12/30/2011 ABN

which is a CON of 12/954,740 11/26/2010 PAT 8153505

which is a CON of 12/720,368 03/09/2010 PAT 7871898

which is a CON of 11/980.664 10/31/2007 PAT 7807549

which is a CON of 10/913,441 08/09/2004 PAT 7387944

which is a CON of 09/505,283 02/16/2000 PAT 6902987

Foreign Applications for which priority is claimed (You may be eligible to benefit from the **Patent Prosecution Highway** program at the USPTO. Please see http://www.uspto.gov for more information.) - None. Foreign application information must be provided in an Application Data Sheet in order to constitute a claim to foreign priority. See 37 CFR 1.55 and 1.76.

Permission to Access Application via Priority Document Exchange: Yes

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Permission to Access Search Results: Yes

Applicant may provide or rescind an authorization for access using Form PTO/SB/39 or Form PTO/SB/69 as appropriate.

If Required, Foreign Filing License Granted: 12/11/2015

The country code and number of your priority application, to be used for filing abroad under the Paris Convention,

is **US 14/957,501**

Projected Publication Date: 03/24/2016

Non-Publication Request: No Early Publication Request: No

Title

METHOD FOR LOW TEMPERATURE BONDING AND BONDED STRUCTURE

Preliminary Class

Statement under 37 CFR 1.55 or 1.78 for AIA (First Inventor to File) Transition Applications: No

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			page 4 of 4				

	Application No.	14/957,501
INFORMATION DISCLOSURE	Filing Date	December 2, 2015
STATEMENT BY APPLICANT	First Named Inventor	Tong et al.
STATEMENT BY APPLICANT	Art Unit	2895
(Multiple sheets used when necessary)	Examiner	Unknown
SHEET 1 OF 5	Attorney Docket No.	TSSRA.012C1

U.S. PATENT DOCUMENTS					
Examiner Initials	Cite No.	Document Number Number - Kind Code (if known) Example: 1,234,567 B1	Publication Date MM-DD-YYYY	Name	Pages, Columns, Lines Where Relevant Passages or Relevant Figures Appear
	1	4,700,466	10-20-1987	Nakagawa et al.	
	2	4,983,251	01-08-1991	Haisma et al.	
	3	5,427,638	06-27-1995	Goetz et al.	
	4	5,451,547	09-19-1995	Himi et al.	

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Examiner Cite Country Code-Number-Kind Code Initials No. Example: JP 1234567 A1		Publication Date MM-DD-YYYY	Name	Pages, Columns, Lines Where Relevant Passages or Relevant Figures Appear	T ¹	
	5	CA 2 399 282 A1	08-23-2001	Ziptronix, Inc.		
	6	JP H08-195334	07-30-1996	Canon Inc.		Т
	7	TW 389965B	05-11-2000	San Fu Chemical Co. Ltd.		Т

	NON PATENT LITERATURE DOCUMENTS				
Examiner Initials	Cite No.	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T ¹		
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Examiner Signature	Date Considered
Examiner Signature	Date Considered

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STATEMENT BY APPLICANT	First Named Inventor	Tong et al.
STATEMENT BY AFFEIGANT	Art Unit	2895
(Multiple sheets used when necessary)	Examiner	Unknown
SHEET 2 OF 5	Attorney Docket No.	TSSRA.012C1

NON PATENT LITERATURE DOCUMENTS				
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STATEMENT BY AFFLICANT	Art Unit	2895
(Multiple sheets used when necessary)	Examiner	Unknown
SHEET 3 OF 5	Attorney Docket No.	TSSRA.012C1

		NON PATENT LITERATURE DOCUMENTS		
Examiner Initials	Litam (haak magazina jaurnal carial aymnacium aatalag ata) data naga(a) yaluma isaua numbar(a) L			
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	Application No.	14/957,501
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STATEMENT BY ALL LICANT	Art Unit	2895
(Multiple sheets used when necessary)	Examiner	Unknown
SHEET 4 OF 5	Attorney Docket No.	TSSRA.012C1

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Examiner Initials	Cite No.	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T ¹
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Examiner Signature Date Considered	
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(19) Canadian Intellectual Property Office

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Un organisme d'Industrie Canada (11) CA 2 399 282

(13) **A1**

- (40) 23.08.2001
- (43) 23.08.2001

(12)

(21) 2 399 282

(51) Int. Cl. 7:

H01L 21/48

(22) 15.02.2001

(85) 05.08.2002

(86) PCT/US01/03683

(87) WO01/061743

(30)

09/505,283 US 16.02.2000

(71)

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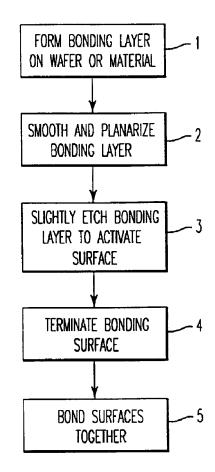
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- (54) PROCEDE DE LIAISON BASSE TEMPERATURE ET STRUCTURE LIEE
- (54) METHOD FOR LOW TEMPERATURE BONDING AND BONDED STRUCTURE

(57)

A method for bonding at low or room temperature includes steps of surface cleaning and activation by cleaning or etching. The method may also include removing by-products of interface polymerization to prevent a reverse polymerization reaction to allow room temperature chemical bonding of materials such as silicon, silicon nitride and SiO2. The surfaces to be bonded are polished to a high degree of smoothness and planarity (2). VSE may use reactive ion etching or wet etching to slighthly etch the surfaces being bonded (3). The surface roughness and planarity are not degraded and may be enhanced by the VSE process. The etched surfaces may be rinsed in solutions such as ammonium hydroxide or ammonium fluoride to promote the formation of desired bonding species on the surfaces (4).



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An agency of Industry Canada CA 2399282 A1 2001/08/23

(21) 2 399 282

(12) DEMANDE DE BREVET CANADIEN CANADIAN PATENT APPLICATION

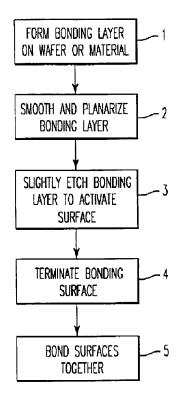
(13) A1

- (86) Date de dépôt PCT/PCT Filing Date: 2001/02/15
- (87) Date publication PCT/PCT Publication Date: 2001/08/23
- (85) Entrée phase nationale/National Entry: 2002/08/05
- (86) N° demande PCT/PCT Application No.: US 2001/003683
- (87) N° publication PCT/PCT Publication No.: 2001/061743
- (30) Priorité/Priority: 2000/02/16 (09/505,283) US

- (51) Cl.Int.⁷/Int.Cl.⁷ H01L 21/48
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(54) Title: METHOD FOR LOW TEMPERATURE BONDING AND BONDED STRUCTURE



(57) Abrégé/Abstract:

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(21) 2 399 282

(13) **A1**

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(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization International Burcau





(43) International Publication Date 23 August 2001 (23.08.2001)

PCT

(10) International Publication Number WO 01/61743 A1

(51) International Patent Classification⁷: H01L 21/48

(21) International Application Number: PCT/US01/03683

(22) International Filing Date: 15 February 2001 (15.02.2001)

(25) Filing Language:

(26) Publication Language:

English

English

(30) Priority Data: 09/505,283

16 February 2000 (16.02.2000) U

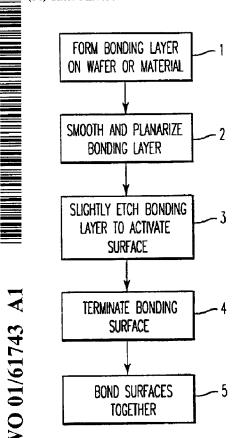
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- (81) Designated States (national): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW.
- (84) Designated States (regional): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European

[Continued on next page]

(54) Title: METHOD FOR LOW TEMPERATURE BONDING AND BONDED STRUCTURE



(57) Abstract: A method for bonding at low or room temperature includes steps of surface cleaning and activation by cleaning or etching. The method may also include removing by-products of interface polymerization to prevent a reverse polymerization reaction to allow room temperature chemical bonding of materials such as silicon, silicon nitride and SiO₂. The surfaces to be bonded are polished to a high degree of smoothness and planarity (2). VSE may use reactive ion etching or wet etching to slighthly etch the surfaces being bonded (3). The surface roughness and planarity are not degraded and may be enhanced by the VSE process. The etched surfaces may be rinsed in solutions such as ammonium hydroxide or ammonium fluoride to promote the formation of desired bonding species on the surfaces (4).

Method For Low Temperature Bonding And Bonded Structure

BACKGROUND OF THE INVENTION

1. Field of the Invention

The present invention relates to bonding of materials at room temperature and, in particular, to bonding of processed semiconductor materials, such as integrated circuit or device substrates, having activated surfaces to achieve high bonding strength adequate for subsequent fabrication and/or a desired application.

2. Background of the Invention

Direct room temperature bonding generally produces weak van der Waals or hydrogen bonding. Annealing is typically required to convert the weak bond to a stronger chemical bond such as a covalent bond. Other wafer bonding techniques including anodic and fusion typically require the application of voltage, pressure and/or annealing at elevated temperature to achieve a sufficient bond strength for subsequent fabrication and/or the desired application. The need to apply voltage, pressure or heat has significantly limited wafer bonding applications because these parameters can damage the materials being wafer bonded, give rise to internal stress and introduce undesirable changes in the devices or materials being bonded. Achieving a strong bond at low temperatures is also critical for bonding of thermally mismatched or thermally sensitive wafers including processed device wafers.

Ultra high vacuum (UHV) bonding is one of the approaches to achieve a low or room temperature strong bond. However, the bonding wafers still have to be pre-annealed at high temperatures, for instance >600° C for silicon and 500° C for GaAs, before cooling down to low or room temperature for bonding. Furthermore, the UHV approach does not generally work on commonly used materials, for example, in SiO₂. It is further also expensive and inefficient.

Adhesive layers can also be used to bond device wafers to a variety of substrates and to transfer device layers at low temperatures. However, thermal and chemical instability, interface bubbles, stress and the inhomogeneous nature of adhesive layers prevent its wide application. It is thus highly desirable to achieve a strong bond at room temperature by bonding wafers in ambient without any adhesive, external pressure or applied electric field.

Low vacuum bonding has been explored as a more convenient alternative to UHV bonding but a bonding energy comparable to the bulk silicon fracture energy using bonded

bare silicon wafer pairs has only be achieved after annealing at $\sim 150^{\circ}$ C. For oxide covered silicon wafer pairs annealing at $\sim 300^{\circ}$ C is required to obtain a high bond energy. It has not been possible to obtain high bonding energies in bonded material using low vacuum bonding at room temperature.

A gas plasma treatment prior to bonding in ambient is known to enhance the bonding energy of bonded silicon pairs at low or room temperature. See, for example, G.L. Sun, Q.-Y. Tong, et al., J. de Physique, 49(C4), 79 (1988); G.G. Goetz, Proc.of 1st Intl. Symp. on Semicond. Wafer Bonding: Science, Technol. and Applications, The Electrochem. Soc., 92-7, 65 (1992); S. Farrens et al., J. Electroch. Soc., 142,3950 (1995) and Amirffeiz et al, Abstracts of 5th Intl. Symp. on Semi. Wafer Bonding: Science, Tech. and Appl., The Electrochemical Society, 99-2, Abstract No. 963 (1999). Although these treatments have increased the bond energy obtainable at low or room temperature, they have only been demonstrated with planar silicon wafers or with silicon wafers using a plasma process that results in oxide being grown on the wafers during the plasma process. Moreover, these treatments have only been used to increase the bond energy by charging or damaging the surface. Furthermore, these treatments have not been used or shown to be applicable to deposited dielectrics or other materials.

Obtaining low or room temperature bonding with a method that is not only applicable to planar silicon and grown oxide surfaces but further to deposited materials and non-planar surfaces with planarized deposited materials will allow generic materials, including processed semiconductor wafers, to be bonded with minimal damage for manufacturing purposes. Such a method based on etching and chemical bonding is described herein.

SUMMARY OF THE INVENTION

It is an object of the invention to provide a method for bonding materials at low or room temperature.

It is another object of the invention to bond materials by cleaning and activating the bonding surfaces to promote chemical bond formation at about room temperature.

It is a further object of the invention to provide a bonding method to bond any solid state material such as processed device or integrated circuit wafers or thermally sensitive or mis-matched materials at or about room temperature.

It is further object of the invention to provide a bonding method to bond processed

device or integrated circuit wafers of different types of devices or different technologies, and transfer a layer of devices or circuits at or about room temperature.

It is another object of the invention to enable a direct wafer bonding method that does not require annealing to achieve a required bond strength.

It is a further object of the invention to provide a method whereby diverse materials including those with non-planar surfaces and deposited materials can be planarized and bonded.

These and other objects are achieved by a method of bonding having steps of forming first and second bonding surfaces, etching the first and second bonding surfaces, and bonding together at room temperature the first and second bonding surfaces after said etching step. The etching may include etching the first and second bonding surfaces such that respective surface roughnesses of the first and second bonding surfaces after said etching are substantially the same as respective surface roughnesses before said etching. The surface roughness may be in a range of 0.1 to 3.0 nm.

The bonding surfaces may be the surface of a deposited insulating material, such as silicon oxide, silicon nitride or a dielectric polymer. The bonding surface may also be the surface of a silicon wafer. Silicon wafers, using either the surface of the wafer or a deposited material on the wafer, may be bonded together. The wafers may have devices or integrated circuits formed therein. The devices and circuits in the wafers bonded together may be interconnected. The wafers may have a non-planar surface or an irregular surface topology upon which a material is deposited to form the bonding surfaces.

Forming at least one of the bonding surfaces may include depositing a polishable material on a non-planar surface. Depositing said polishable material may include depositing one of silicon oxide, silicon nitride or a dielectric polymer. The bonding surfaces may be polished using a method such as chemical-mechanical polishing. The surfaces may also be etched prior to the polishing.

The etching step may also include activating the first and second bonding surfaces and forming selected bonding groups on the first and second bonding surfaces. Bonding groups may also be formed capable of forming chemical bonds at approximately room temperature, and chemical bonds may be formed between the bonding surfaces allowing bonded groups to diffuse or dissociate away from an interface of the bonding surfaces. The chemical bonds can

increase the bonding strength between the bonding surfaces by diffusing or dissociating away said bonding groups.

After said etching step, the bonding surfaces may be immersed in a solution to form bonding surfaces terminated with desired species. The species may comprise at least one of a silanol group, an NH₂ group, a fluorine group and an HF group. Also, a monolayer of one of a desired atom and a desired molecule may be formed on the bonding surface. Terminating the surface may include rinsing said bonding materials in an ammonia-based solution after said slightly etching. The ammonia-based solution may be ammonium hydroxide or ammonium fluoride.

The method may also include exposing the bonding surfaces to one of an oxygen, argon, NH₃ and CF₄ RIE plasma process. Silicon dioxide may be deposited as to form the bonding surfaces, and etched using the RIE process.

The etching process may create a defective or damaged zone proximate to the bonding surfaces. The defective or damaged zone can facilitate the removal of bonding by-products through diffusion or dissociation.

The method may also include steps of forming first and second bonding surfaces, etching the bonding surfaces, terminating the bonding surfaces with a species allowing formation of chemical bonds at about room temperature, and bonding the bonding surfaces at about room temperature, or may include steps of forming the bonding surfaces each having a surface roughness in a range of 0.1 to 3 nm, removing material from the bonding surfaces while maintaining said surface roughness, and directly bonding the bonding surfaces at room temperature with a bonding strength of at least 500 mJ/m², at least 1000 mJ/m², or at least 2000 mJ/m².

The objects of the invention may also be achieved by a bonded device having a first material having a first etched bonding surface, and a second material having a second etched bonding surface directly bonded to the first bonding surface at room temperature having a bonding strength of at least 500 to 2000 mJ/m². The bonding surfaces may be being activated and terminated with a desired bonding species, and the desired species may include a monolayer of one of a desired atom and a desired molecule on said bonding surface or at least one of a silanol group, an NH₂ group, a fluorine group and an HF group. The bonding surfaces may each have a defective region located proximate to said first and second bonding

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surfaces, respectively.

The first material may include a surface of a first semiconductor wafer having devices formed therein, and the second material may include a surface of a second semiconductor wafer having devices formed therein. Devices in the wafers may be interconnected, and the wafers may be of different technologies. The wafers may also have an integrated circuit formed therein, and devices or circuits in the wafers may be interconnected.

One of said first and second wafers may be a device region after removing a substantial portion of a substrate of said one of said first and second wafers. The wafers may have an irregular surface topology.

The first material may include a first wafer containing electrical devices and having a first non-planar surface, and the first bonding surface may include a polished and etched deposited oxide layer on said first non-planar surface. The second material may include a second wafer containing electrical devices and having a second non-planar surface, and the second bonding surface may include a polished, planarized and slightly etched deposited oxide layer on the second non-planar surface.

The first material may include a first wafer containing electrical devices and having a first surface with irregular topology, and the first bonding surface may include a polished, planarized and slightly etched deposited oxide layer on the first surface. The second material may include a second wafer containing electrical devices and having a second surface with irregular topology, and the second bonding surface may include a polished, planarized and slightly etched deposited oxide layer on the second surface.

The bonded device according to the invention may also include a first material having a first etched and activated bonding surface terminated with a first desired bonding species, and a second material having a second etched and activated bonding surface terminated with a second desired bonding species bonded to the first bonding surface at room temperature.

BRIEF DESCRIPTION OF THE DRAWINGS

A more complete appreciation of the invention and many of the attendant advantages thereof are readily obtained as the same become better understood by reference to the following detailed description when considered in connection with the accompanying drawings, wherein:

- FIG. 1 is a flow chart of the method according to the invention;
- FIG. 2 is a flow chart of an example of the method according to the invention;
- FIGS. 3A-3E are diagrams illustrating a first embodiment of a method according to the invention;
- FIG. 4 is a diagram illustrating bonding according to the invention using silicon oxide;
 - FIG. 5 is a diagram illustrating bonding according to the invention using silicon;
 - FIGS. 6A and 6B are graphs of room temperature bonding energy versus storage time;
 - FIG. 7 is a diagram of a bonding fixture used in the invention; nd
- FIG. 8 is a fluorine concentration profile by SIMS (Secondary Ion Mass Spectroscopy) near the bonding interface of deposited oxide covered silicon wafers that were very slight etched by diluted HF before bonding.

DESCRIPTION OF THE PREFERRED EMBODIMENTS

Referring to FIGS. 1 and 3A-3E, a first embodiment of the method according to the invention will be described. Wafer 30, preferably a processed semiconductor device wafer and more preferably a processed silicon device wafer, contains a device layer 31 with processed devices. Device layer 31 may contain a number of layers and include surface regions of wafer 30. The surface topography of layer 31 is typically nonplanar. Layer 31 may also represent a processed integrated circuit containing any number of layers such as active devices, interconnection, insulation, etc.

The integrated circuit may be fully processed, or partially processed where the remaining processing is performed after the bonding process. The processing after the bonding may include full or partial substrate removal or via formation between the bonded wafers for interconnection.

On layer 31 a bonding layer 32 is formed (step 1, FIG. 1). Bonding layer 32 may be any solid state material or mixed materials which can be deposited or formed at low temperatures and can be polished to a sufficiently smooth surface. Layer 32 may be an insulator, such as SiO₂, silicon nitride, amorphous silicon formed using chemical vapor deposition (CVD) or plasma-enhanced CVD (PECVD), sputtering or by evaporation. Other materials such as polymers, semiconductors or sintered materials may also be used. Layer 32

should have thickness greater than the surface topography of layer 31.

The surface 33 of layer 32 is planarized and smoothed, as shown in step 2 of FIG. 1 and in FIG. 3B. It is noted that the roughness/planarity of surface 33 is exaggerated in FIG. 3A for illustrative purposes. This step may be accomplished using chemical-mechanical polishing. Surface 33 is preferably polished to a roughness of about no more than about 3 nm and preferably no more than about 0.1 nm and be substantially planar. The surface roughness values are typically given as root-mean square (RMS) values. Also, the surface roughness may be given as mean values which are nearly the same as the RMS values. After polishing surface 33 is cleaned and dried to remove any residue from the polishing step. Polished surface 33 is preferably then rinsed with a solution.

The bonding surface may also be etched prior to polishing to improve the planarity and/or surface roughness. The etching can be effective to remove high spots on the bonding surface by selective etching of the high spots using, for example, standard photolithographic techniques. For example, a layer of silicon nitride can be embedded within a silicon dioxide bonding layer 32 that can serve as an etch stop when using a solution containing HF. The etch stop material may be used to improve uniformity, reproducibility, and manufacturability.

FIG. 3B illustrates layer 32 having upper surface 34 after the polishing/planarization and cleaning steps. Surface 34 then undergoes an activation process (step 3, FIG. 1). This activation process is an etching process and preferably a very slight etch (VSE) process. The term VSE means that the root-mean-square micro-roughness (RMS) of the very slightly etched surface remains at approximately the unetched value, typically < 0.5 nm and preferably in the range of 0.1 nm to 3 nm. The optimum amount of material removed depends upon the material and the method used for removal. Typical amounts removed vary from Angstroms to a few nanometers. It is also possible to remove more material. VSE also includes the breaking of bonds on the treated surfaces and can occur without significant removal of material. The VSE is distinct from simple modification of the surface by, for example, charging the surface with electronic charge or damaging the surface layer. In a first example of the method according to the invention, the VSE process consists of a gas or mixed gas (such as oxygen, argon, nitrogen, CF4, NH3) plasma process at a specified power level for a specified time (FIG. 3C). The power and duration of the plasma process will vary depending upon the materials used to obtain the desired bond energy. Examples are given

below, but in general, the power and duration will be determined empirically.

The plasma process may be conducted in different modes. Both reactive ion etch (RIE) and plasma modes may be used, as well as an inductively-coupled plasma mode (ICP). Sputtering may also be used. Data and examples are given below in both the RIE and plasma modes.

The VSE process etches the surface very slightly via physical sputtering and/or chemical reaction and preferably is controlled to not degrade the surface roughness of the bonding surfaces. The surface roughness may even be improved depending upon the VSE and materials etched. Almost any gas or gas mixture that will not etch surface 34 excessively can be used for the room temperature bonding method according to the invention.

The VSE serves to clean the surface and break bonds of the oxide on the wafer surface. The VSE process can thus enhance the surface activation significantly. A desired bonding species can be used to terminated on surface 34 during the VSE by proper design of the VSE. Alternatively, a post-VSE treatment that activates and terminates the surface with a desired terminating species during the post-VSE process may be used.

The desired species further preferably forms a temporary bond to the surface 34 atomic layer, effectively terminating the atomic layer, until a subsequent time that this surface can be brought together with a surface terminated by the same or another bonding species 36 as shown in FIG. 3D. Desired species on the surfaces will further preferably react with each other when they are in sufficiently close proximity allowing chemical bonding between surfaces 34 and 36 at low or room temperature that is enhanced by diffusion or dissociation and diffusion of the reacted desired species away from the bonding interface.

The post-VSE process preferably consists of immersion in a solution containing a selected chemical to generate surface reactions that result in terminating the bonding surface 34 with desired species. The immersion is preferably performed immediately after the VSE process. The post-VSE process may be performed in the same apparatus in which the VSE process is conducted. This is done most readily if both VSE and post-VSE processes are either dry, i.e, plasma, RIE, ICP, sputtering, etc, or wet, i.e., solution immersion. A desired species preferably consists of a monolayer or a few monolayers of atoms or molecules.

The post-VSE process may also consist of a plasma, RIE, or other dry process whereby appropriate gas chemistries are introduced to result in termination of the surface

with the desired species. The post-VSE process may also be a second VSE process. The termination process may also include a cleaning process where surface contaminants are removed without VSE. In this case, a post-cleaning process similar to the post-VSE processes described above then results in a desired surface termination.

The post-VSE or post-cleaning process may or may not be needed to terminate surfaces with desired species if the activated surface bonds by the cleaning or VSE process are subsequently sufficiently weakly surface reconstructed and can remain sufficiently clean before bonding such that subsequent bonding with a similar surface can form a chemical bond.

The wafers are optionally rinsed then dried. Two wafers are bonded by aligning them (if necessary) and bringing them together to form a bonding interface. As shown in FIG. 3D, a second wafer 35 has been processed in the manner shown in FIG. 3C to prepare bonding surface 36. The two wafers are brought together by, for example, commercially available wafer bonding equipment (not shown) to initiate bonding interface 37 (FIG. 3E).

A spontaneous bond then typically occurs at some location in the bonding interface and propagates across the wafer. As the initial bond begins to propagate, a chemical reaction such as polymerization that results in chemical bonds takes place between species used to terminate surfaces 34 and 36 when the surfaces are in sufficient proximity. The bonding energy is defined as the specific surface energy of one of the separated surfaces at the bonding interface that is partially debonded by inserting a wedge. The by-products of the reaction then diffuse away from the bonding interface to the wafer edge or are absorbed by the wafers, typically in the surrounding materials. The by-products may also be converted to other by-products that diffuse away or are absorbed by the wafers. The amount of covalent and/or ionic bonding may be increased by removal of converted species resulting in further increase in bond strength.

FIGS. 4A-4E show surface conditions and the bonding propagation to form covalent bonds in a the case of a planar Si wafer covered with silicon oxide. On Si wafer 40 an SiO₂ layer 41 is formed, which has been polished and planarized. Surface 42 of layer 41 is subjected to the VSE process to produce an activated surface (FIG. 4A). On a second wafer 44 a second SiO₂ layer 45 is formed, and surface 46 is subjected to a VSE process to activate surface 46 (FIG. 4B). Desired species are terminated on surface 46 and are shown as lines 43

in FIG. 4C. Either or both of a VSE and post-VSE processes are used to properly terminate surface 46. While not shown, surface 42 may also be terminated using a post-VSE process. Wafer 44 is brought together with wafer 40 (FIG. 4D) and bonds 46 begin to form. the bonding propagates and by-products are removed (indicated as arrows 47) and chemical bonds (such as covalent) are formed, as shown in FIG. 4E.

The bonding immediately after the RIE process may use a special bonding fixture allowing immediate in situ bonding of the etched wafers. A diagram of the fixture is shown in FIG. 7. In plasma chamber 75 are two wafers to be bonded 70 disposed on RF electrodes 76 and 77. A plasma is formed in zone 79 by the application of RF power to the electrodes via moveable vacuum RF power feedthrough 74 and by the introduction of an appropriate gas or gas mixture through gas feedthrough 73. Element 71 is a vacuum feedthrough for mechanical actuator (not shown) to retract retractable spacer 72. Chamber 75 is pumped down to a desired vacuum level via pumps (not shown) and chamber inlet 78. In the case where a post-VSE process or post cleaning process is also a dry process, as discussed above, the VSE and post-VSE or post-cleaning may be conducted in chamber 75.

After the plasma treatment to conduct the VSE process, the mechanical spacers 72 are retracted by the mechanical actuator and the wafers 70 are moved into contact with to begin the bonding process. The bonded wafers are then moved from the chamber into ambient or into another vacuum chamber (not shown) and stored for a desired period to allow the bonding to propagate by a wafer handling system (not shown).

The materials of the bonding layers preferably have an open structure so that the byproducts of the polymerization reaction can be easily removed. The bonding species on the
opposing bonding surfaces must be able to react at room temperature to form a strong or
chemical bond. The bond energy is sufficiently high to virtually eliminate slippage between
wafers after subsequent heat treatments associated with a subsequent processing or operation
when wafers have different thermal expansion coefficients. Lack of slippage is manifest by a
lack of wafer bowing upon inspection after the subsequent processing or operation.

In order to achieve the high bonding energies, it is preferable for at least one of the wafers to be as thin as possible because a thin wafer allows compliance to accommodate a lack of perfect surface planarization and smoothness. Thinning to thickness of about 10 mils to 10 microns is effective.

The bonded wafers are preferably stored at ambient or at low or room temperature after bonding to allow removal of species or converted species for a specified period of time depending upon the materials and species used. Twenty four hours is usually preferable. The storage time is dependent upon the type of plasma process used. Chemical bonds may be obtained more quickly, in a matter of minutes, when certain plasma processes such as an Ar plasma are used. For example, 585 mJ/m² bonds were obtained in immediately after bonding and over 800 mJ/m² were observed after 8 hours for deposited oxides etched by an Ar plasma followed by NH₄OH dip.

Annealing the bonded wafers during bonding may increase the bonding strength. The annealing temperature should be below 200° C and may be typically in the range of 75-100° C. Storing the bonded wafers under vacuum may facilitate the removal of residual gasses from the bonding surfaces, but is not always necessary.

All of the processes above may be carried out at or near room temperature. The wafers are bonded with sufficient strength to allow subsequent processing operations (lapping, polishing, substrate removal, chemical etching, lithography, masking, etc.).

Bonding energies of approximately 500-2000 mJ/m² or more can be achieved (see FIG. 6A).

At this point (FIG. 3E) it is possible to remove a part or all of the substrate of wafer 35 by, for instance, lapping and etch back. The layer of devices of wafer 35 is thus transferred onto wafer 30. The devices from the two layers may be interconnected. Additional device or circuit layers may be bonded and interconnected to form a multilayer structure. Different types of wafers, devices or circuits may be bonded, as well as different technologies (i.e. CMOS and bipolar or III-V HBT and Si CMOS). Other elements or materials such as thermal spreaders, surrogate substrates, antennas, wiring layers, a preformed multi-layer interconnects, etc. may be bonded to produce different types of circuits or systems, as desired.

In an example, shown in FIG.2, PECVD SiO₂ is deposited on a Si wafer containing devices. Surface 34, after the plasma (such as argon, oxygen or CF₄) treatment, is mainly terminated by Si-OH groups due to the availability of moisture in the plasma system and in air. After the plasma treatment, the wafers are immediately immersed in solution such as ammonium hydroxide (NH₄OH), NH₄F or HF for a period such as between 10 and 120 seconds. After immersing the wafers in the NH₄OH solution, many Si-OH groups are

replaced by Si-NH2 groups according to the following substitution reaction:

$$2Si-OH + 2NH4OH - 2Si-NH2 + 4HOH$$
 (1)

Alternatively, many Si-F groups are terminating on the PECVD SiO_2 surface after an NH_4F or HF immersion.

The hydrogen bonded Si-NH2:Si-OH groups or Si-NH2:Si-NH2 groups across the bonding surfaces can polymerize at room temperature in forming Si-O-Si or Si-N-N-Si (or Si-N-Si) covalent bonds:

$$Si-NH2 + Si-OH \rightarrow Si-O-Si + NH3$$
 (2)

$$Si-NH2 + Si-NH2 \rightarrow Si-N-N-Si + 2H2$$
 (3)

Alternatively, the HF or NH₄F dipped oxide surfaces are terminated by Si-F groups in addition to Si-OH groups. Since HF or NH₄F solution etches silicon oxide strongly, their concentrations must be controlled to an adequately low level, and the immersion time must be sufficiently short. This is an example of a post-VSE process being a second VSE process. The covalent bonds across the bonding interface are formed due to the polymerization reaction between hydrogen bonded Si-HF or Si-OH groups:

$$Si-HF + Si-HF \rightarrow Si-F-F-Si + H2$$
 (4)

$$Si-F + Si-OH \rightarrow Si-O-Si + HF$$
 (5)

FIG. 8 shows the fluorine concentration profile of bonded thermal oxide covered silicon wafers that were dipped in 0.05% HF before room temperature bonding. A fluorine concentration peak is clearly seen at the bonding interface. This provides evidence of the chemical process described above where the desired species are located at the bonding interface.

Since reaction (2) is reversible only at relatively high temperatures of ~500° C, the formed siloxane bonds should not be attacked by NH₃ at lower temperatures. It is known that H₂ molecules are small and diffuse about 50 times quicker than water molecules in oxide. The existence of a damaged layer near the surface of an adequate thickness i.e. a few nm, will facilitate the diffusion or dissolution of NH₃, and HF and hydrogen in reactions (2), (3), (4) and/or (5) in this layer and enhancement of the chemical bond. The three reactions result in a higher bonding energy of SiO₂/SiO₂ bonded pairs at room temperature after a period of storage time to allow NH₃ or H₂ to diffuse away.

In the example of FIG. 2, the plasma treatment may create a damaged or defective

area in the oxide layer near the bonding surface. The zone extends for a few monolayers. The damaged or defective area aids in the removal of bonding by-products. Efficient removal of the bonding by-products improves the bonding strength since the by-products can interfere with the bonding process by preventing high-strength bond from forming..

Many different surfaces of materials may be smoothed and/or planarized, followed by a cleaning process, to prepare for bonding according to the invention. These materials can be room temperature bonded by mating surfaces with sufficient planarity, surface smoothness, and passivation that includes cleaning, and/or VSE, activation and termination. Amorphous and sintered materials, non-planar integrated circuits, and silicon wafers are examples of such materials. Single crystalline semiconductor or insulating surfaces, such as SiO₂ or Si surfaces, can also be provided with the desired surface roughness, planarity and cleanliness. Keeping the surfaces in high or ultra-high vacuum simplifies obtaining surfaces sufficiently free of contamination and atomic reconstruction to achieve the strong bonding according to the invention. Other semiconductor or insulator materials such as InP, GaAs, SiC, sapphire, etc., may also be used. Also, since PECVD SiO₂ may be deposited on many types of materials at low temperatures, many different combinations of materials may be bonded according to the invention at room temperature. Other materials may also be deposited as long as appropriate processes and chemical reactions are available for the VSE, surface activation, and termination.

For example, the method may also be used with silicon nitride as the bonding material. Silicon nitride may be bonded to silicon nitride, or to silicon dioxide and silicon. Silicon oxide may also be bonded to silicon. Other types of dielectric materials may be bonded together including aluminum nitride and diamond-like carbon.

The method may be applied to planar wafers having no devices or circuits and one wafer with devices and circuits. The planar wafer may be coated with a bonding layer, such as PECVD oxide or amorphous silicon, and then processed as described above to bond the two wafers. The planar wafer may not need to be coated with a bonding layer if it has sufficient smoothness and planarity and the proper bonding material.

As can be appreciated, the bonding process may be repeated with any number of wafers, materials or functional elements. For example, two device or IC wafers may be joined, followed by removing one of the exposed substrates to transfer a layer or more of

devices, or just the active regions of an IC.

The bonding according to the invention may be applied to joining different types of materials. For example, a silicon wafer can be bonded to another silicon wafer, or bond to an oxidized silicon wafer. The bare silicon wafer and the oxide covered wafer are immersed in HF, NH₄F and/or NH₄OH and bonded after drying. The time for the immersion should be less than about twenty minutes for the silicon wafer covered with the thin oxide since the NH₄OH solution etches silicon oxide. Since HF and NH₄F etches oxides strongly, very diluted solutions, preferably in 0.01-0.2% range should be used for dipping of the silicon wafers.

After drying the silicon wafer and the oxide-covered wafer are bonded in ambient at room temperature. Reactions (2), (3), (4) and/or (5) take place at the bonding interface between the two wafers. The plasma-treated wafers may also be immersed in deionized water instead of the NH₄OH solution.

The silicon bonding may be conducted with a bare silicon wafer, i.e. having a native oxide or a silicon wafer having an oxide layer formed on its surface as described above. During the oxygen plasma treatment, the native oxide which if formed on the bare silicon wafer is sputter etched, and the oxide layer formed on the silicon surface is etched. The final surface is an activated (native or formed) oxide. When rinsed in deionized water, the activated oxide surface is mainly terminated with Si-OH groups. Since oxide growth in oxygen plasma has been found to have less water than in normal native oxide layers, the water from the original bonding bridge and generated by the following polymerization reaction (6) can be absorbed into the plasma oxide readily.

$$Si-OH + Si-OH \rightarrow Si-O-Si + H2O$$
 (6)

FIGS. 5A-5E illustrate bonding two silicon wafers. Wafers 50 and 52 have respective surfaces 51 and 53 with native oxides (not shown) subjected to a VSE process. Surface 53 is in FIG. 5C is shown terminated with a desired species 54. The two wafers are brought together and bonds 55 begin to form (FIG. 5D). The bonding propagates and bonding byproducts, in this case H₂ gas, are removed. The by-products being removed are shown as arrows 56 in FIG. 5E.

In addition to removal of the water from the bonding interface by dissolving into the plasma activated oxide of the oxidized silicon wafer, the water can also diffuse through the

thin oxide layer on the bare silicon wafer to react with silicon. As the silicon surface underneath the oxide has a damaged or defective zone, extending for a few monolayers, the water molecules that diffuse through the oxide layer and reach the damaged or defective zone can be converted to hydrogen at room temperature and be removed readily:

$$Si + 2H_2O - SiO_2 + 2H_2 \tag{7}$$

The reverse reaction of (6) is thus avoided and the room temperature bonding energy increases enormously due to the formation of covalent Si-O-Si bonds.

If a relatively thick (~5 nm) oxide layer is formed, it will take a long period of time for the water molecules to diffuse through this thick layer. On the other hand, if after the plasma treatment a thin oxide layer is left or a too narrow defective zone is formed, water that can reach the silicon surface may not react sufficiently with the silicon and convert to hydrogen. In both cases the bonding energy enhancement will be limited. The preferred oxygen plasma treatment thus leaves a minimum plasma oxide thickness (e.g., about 0.1-1.0 nm) and a reasonably thick defective zone (e.g., about 0.1-0.3 nm) on the silicon surface.

In a second embodiment, the VSE process uses wet chemicals. For example, an InP wafer having a deposited silicon oxide layer, as in the first embodiment, and a device layer are bonded to a AlN substrate having a deposited oxide layer. After smoothing and planarizing the InP wafer bonding surface and the AlN wafer bonding surface, both wafers are cleaned in an standard RCA cleaning solution. The wafers are very slightly etched using a dilute HF aqueous solution with an HF concentration preferably in the range of 0.01 to 0.2%. About a few tenths of a nm is removed and the surface smoothness is not degraded as determined by AFM (atomic force microscope) measurements. Without deionized water rinse, the wafers are spin dried and bonded in ambient air at room temperature. The resulting bonding energy has been measured to reach ~700 mJ/m² after storage in air. After annealing this bonded pair at 75°C the bonding energy of 1500 mJ/m² was obtained., The bonding energy has been measured to reach silicon bulk fracture energy (about 2500 mJ/m²) after annealing at 100°C. If the wafers are rinsed with deionized water after the HF dip, the bonding energy at 100°C is reduced to 200 mJ/m², that is about one tenth of that obtained without the rinse. This illustrates the preference of F to OH as a terminating species.

In a third embodiment the VSE process consists of 0.1% HF etching followed by 5 min dip in 0.02% HN₄F solution of thermally oxidized silicon wafers at room temperature

after a standard cleaning process. Without rinsing in deionized water, the wafers are bonded after spin drying at room temperature. The bonding energy of the bonded pairs reaches ~1700 mJ/m² after 100°C annealing. If the wafers are rinsed in de-ionized water after the HF etching before bonding, the bonding energy of bonded pairs is only 400 mJ/m², again illustrating the preference of F to OH as a terminating species.

Dilute NH₄F is used in the VSE process to etch silicon oxide covered wafers in a fourth embodiment. The concentration of the NH₄F should be below 0.02% to obtain the desired bonding. The bonding energy of ~600 mJ/m² can be achieved at room temperature after storage.

A fifth embodiment of the invention is used to bond Si surfaces having a native oxide of about 1 nm in thickness. In the fifth embodiment, after cleaning the Si surface by a standard RCA1 cleaning process, a VSE process using 5 min etching in 70% HNO₃ + diluted HF (preferably 0.01 to 0.02%) is performed. Wafers are pulled out of the solution vertically with a basically hydrophobic surface. Without rinsing in water, the wafers were bonded at room temperature in air. In this process covalent bonding occurs at room temperature with measured bonding energies typically about 600 mJ/m². This bonding energy is significantly increased to 1300 mJ/m² after annealing at 75°C and reaches the fracture energy of bulk silicon (about 2500 mJ/m²) at a temperature of 100° C.

Instead of 70% HNO₃, diluted HNO₃ with water can be used in the solution to achieve similar results. According to AMF measurements and high resolution transmission electron microscopy measurement results, the silicon is etched in the dilute HNO₃ VSE process at a rate of 0.1-0.15 nm/min and a new thick oxide 2.5-3.5 nm in thickness is formed.

As further embodiments, the VSE process may consist of a dry etch that has chemical and/or physical components. For a bare Si surface, chemical etching may result from SF₄/H₂ gas mixture while physical etching may result from Ar etch. For a silicon oxide surface, chemical etching may use CF₄ while physical etching may use oxygen or argon gas. It is also possible to use a thermally stable polymer material for the bonding materials and bond two polymer surfaces together. Examples are polyimides or spin-on materials.

The mechanisms governing the increased bond energy at low or room temperature are similar. A very slight etching (VSE) of the bonding wafers by plasma to clean and activate the surfaces, and improve removal of by-products of interface polymerization to prevent the

undesirable reverse reaction and rinse in appropriate solution to terminate the surface with desired species to facilitate room temperature covalent bonding. The oxide covered wafer bonding case is similar except that a different surface termination is preferred. In bare silicon wafer bonding, the highly reactive surface layers of oxide and silicon to allow water adsorption and conversion to hydrogen should be formed. The highly reactive layers can be a plasma thin oxide layer and a damaged silicon surface layer. The oxide on the silicon wafer will also have some damage. Not only O_2 plasma but also plasma of other gases (such as Ar, CF_4) are adequate. Because during and after VSE the silicon surface is readily to react with moisture to form an oxide layer, and the underlying damaged silicon layer is created by VSE. Since the VSE and by-products removal methods are rather general in nature, this approach can be implemented by many means and apply to many materials.

EXAMPLE 1

In a first example, three inch <100>, 1-10 ohm-cm, boron doped silicon wafers were used. PECVD oxide was deposited on some of the silicon wafers. For comparison, thermal oxidized silicon wafers were also studied. The PECVD oxide thickness was 0.5 µm and 0.3 µm on the front side and the back side of the wafers, respectively. Oxide is deposited on both sides of the wafer to minimize wafer bow during polishing and improve planarization. A soft polish was performed to remove about 30 nm of the oxide and to smooth the front oxide surface originally having a root mean square of the micro-roughness (RMS) of ~0.56 nm to a final ~0.18 nm. A modified RCA1 solution was used to clean the wafer surfaces followed by spin-drying.

Two wafers were loaded into the plasma system, both wafers are placed on the RF electrode and treated in plasma in RIE mode. For comparison, some wafers were treated in plasma mode in which the wafers were put on the grounded electrode. An oxygen plasma was used with a nominal flow rate of 16 scc/m. The RF power was 20-400 W (typically 80 W) at 13.56 MHz and the vacuum level was 100 mTorr. The oxide covered wafers were treated in plasma for times between 15 seconds to 5 minutes. The plasma treated silicon wafers were then dipped in an appropriate solution or rinse with de-ionized water followed by spin-drying and room temperature bonding in air. Some of the plasma treated wafers were also directly bonded in air without rinse or dipping.

The bonding energy was measured by inserting a wedge into the interface to measure the crack length according to the equation:

$$\gamma = \frac{3t_b^2 E_1 t_{wl}^3 E_2 t_{rw2}^3}{16L^4 \left(E_1 t_{wl}^3 + E_2 t_{w2}^3 \right)}$$

E and tw are the Young's modulus and thickness for wafers one and two and the is the thickness of a wedge inserted between the two wafers that results in a wafer separation of length L from the edge of the wafers.

The room temperature bonding energy as a function of storage time of bonded plasma treated oxide covered silicon wafers is shown in FIG. 6A. This figure shows measured room temperature bonding energy versus storage time for 4 different cases as shown. The results can be summarized as follows: (1) for dipped and bonded RIE plasma treated oxide wafers, the room temperature bonding energy increases with storage time and reaches a stable value after ~20 h in air or at low vacuum; (2) RIE mode results in higher bonding energies than plasma mode; (3) too short a plasma exposure time or too low a plasma power provides a small or negligible increase in bond energy; (4) NH₄OH dip after plasma treatment shows a much higher increase in bonding energy than water rinse; (5) direct bonding in air after plasma treatment without dipping or rinse shows an almost constant bonding energy with time. The bonding energy of the directly bonded wafer pairs immediately after room temperature bonding is slightly higher than the de-ionized water rinsed or NH₄OH dipped wafer pairs.

FIG. 6B shows room temperature bonding of Si and AlN wafers with PECVD oxide deposited layers. After about 100 h of storage time a bonding energy of over 2000 mJ/m² were observed.

Comparing different bonding materials, the bonding energy as a function of storage time of O_2 plasma treated thermally oxidized silicon wafer pairs is similar to wafers with PECVD oxide, although the values of the room temperature bonding energy are somewhat lower.

After ~24 h storage in air at room temperature, the bonding energy as high as ~1000 mJ/m2 was reached in the RIE mode plasma treated and NH₄OH dipped PECVD oxide

covered wafer pairs. Since the maximum bonding energy of a van der Waals bonded silicon oxide covered wafer pairs is about 200 mJ/m2, a large portion of the bonding energy is attributed to the formation of covalent bonds at the bonding interface at room temperature according to the above equation.

EXAMPLES 2-3

The above process was applied to bond processed InP wafers (600 µm thick) to AlN wafers (380 µm thick), or to bond processed Si (380 µm thick) and InP (600 µm thick) wafers, as second and third examples. The processed InP device wafers are covered with PECVD oxide and planarized and smoothed by chemical-mechanical polishing CMP. A PECVD oxide layer is also deposited on the AlN wafers and is planarized and smoothed to improve the RMS surface roughness. The processed Si and processed InP wafers are deposited with PECVD oxide and planarized and smoothed using CMP. After VSE similar to the example 1 bonding at room temperature, the bonded wafers are left in ambient air at room temperature.

After 24 hours storage at room temperature, bonding energy of 1000 mJ/m2 and 1100 mJ/m2 were achieved for the InP/Si and InP/AlN bonded pairs, respectively. For processed Si (380 µm thick) /oxide covered AlN (280 µm thick) wafer pairs, the bonding energy at room temperature as high as 2500 mJ/m2 has been achieved. These room temperature bonded plasma treated wafer pairs have sufficient bonding strength to sustain subsequent substrate lapping and etching and other typical semiconductor fabrication processes before or after substrate removal.

The InP substrate in the room temperature bonded InP/AlN pairs was lapped with 1900# Al2O3 powder from initial 600 μ m thick to ~50 μ m thick followed by etching in an HCl/H3PO4 solution to leave about a 2.0 μ m thick InP device layer on the AlN or Si wafer. The water and etching solution did not penetrate into the bonding interface.

Surfaces are sputter etched by energetic particles such as radicals, ions, photons and electrons in the plasma or RIE mode. For example, the O₂ plasma under conditions that bring about the desired VSE is sputter-etching about 2 Å/min of PECVD oxide as measured by a reflectance spectrometry. For thermal oxide the sputter etching rate is about 0.5 Å/min. The thickness of oxide before and after plasma treatment was measured by a reflectance

spectrometry and averaged from 98 measured points on each wafer. The etching by O_2 plasma has not only cleaned the surface by oxidation and sputtering but also broken bonds of the oxide on the wafer surfaces.

However, the surface roughness of plasma treated oxide surfaces must not be degraded by the etching process. AFM measurements show that compared with the initial surface roughness, the RMS of the O_2 plasma treated oxide wafers was \sim 2 Å and did not change noticeably. On the other hand, if the etching is not sufficiently strong, the bonding energy enhancement effect is also small. Keeping other conditions unchanged when the O_2 plasma treatment was performed with plasma mode rather than RIE mode, the etching of oxide surfaces is negligible and the oxide thickness does not change. The final room temperature bonding energy is only 385 mJ/m2 compared to 1000 mJ/m2 of RIE treated wafers (see FIG. 6A).

Other gas plasma has shown a similar effect. CF_4/O_2 RIE was used to remove ~4 nm of PECVD oxide from the wafer surfaces prior to bonding. The bonding energy of room temperature bonded PECVD oxide covered silicon wafers was also enhanced significantly in this manner and exceeds 1000 mJ/m2 after sufficient storage time (see also FIG. 6A).

An argon plasma has also been used for the VSE with a nominal flow rate of 16 scc/m. The RF power was typically 60 W at 13.56 MHz and the vacuum level was 100 mTorr. The oxide covered silicon wafers were treated in plasma in RIE mode for times between 30 seconds to 2 minutes. The plasma treated silicon wafers were then dipped in an NH4OH solution followed by spin-drying and room temperature bonding in air. The bonding energy reached \sim 800 mJ/m² at room temperature after only 8 h storage in air.

Obviously, numerous modifications and variations of the present invention are possible in light of the above teachings. It is therefore to be understood that within the scope of the appended claims, the invention may be practiced otherwise than as specifically described herein.

Claims:

1. A bonding method, comprising:

forming first and second bonding surfaces;

etching said first and second bonding surfaces; and

bonding together at room temperature said first and second bonding surfaces after said etching step.

2. A method as recited in claim 1, wherein said etching step comprises:

etching said first and second bonding surfaces such that respective surface roughnesses of said first and second bonding surfaces after said etching are substantially the same as respective surface roughnesses before said etching.

3. A method as recited in claim 2, comprising:

forming said first and second bonding surfaces to have a surface roughness in a range of 0.1 to 3.0 nm.

- 4. A method as recited in claim 1, wherein said etching step comprises:
- activating said first and second bonding surfaces and forming selected bonding groups on said first and second bonding surfaces.
 - 5. A method as recited in claim 4, comprising:

forming bonding groups capable of forming chemical bonds at approximately room temperature.

6. A method as recited in claim 1, comprising:

forming chemical bonds between said bonding surfaces allowing bonded groups to diffuse or dissociate away from an interface of said first and second bonding surfaces.

7. A method as recited in claim 6, comprising:

increasing bonding strength between said first and second bonding surfaces by diffusing or dissociating away said bonding groups.

- 8. A method as recited in claim 1, wherein said etching step comprises:
- forming a monolayer of one of a desired atom and a desired molecule on said bonding surface.
 - 9. A method as recited in claim 8, wherein said etching step comprises:

forming a few monolayers of one of a desired atom and a desired molecule on said bonding surface.

10. A method as recited in claim 1, comprising:

after said etching step, immersing said first and second bonding surfaces in a solution to form bonding surfaces terminated with desired species.

- 11. A method as recited in claim 10, wherein said species comprise at least one of a silanol group, an NH₂ group, a fluorine group and an HF group.
 - 12. A method as recited in claim 10, wherein said etching step comprises:

forming a monolayer of one of a desired atom and a desired molecule on said bonding surface.

- 13. A method as recited in claim 1, wherein said etching comprises: exposing said first and second bonding surfaces to a plasma.
- 14. A method as recited in claim 1, comprising:

exposing said first and second bonding surfaces to one of an oxygen, argon, NH₃ and CF₄ plasma process.

15. A method as recited in claim 14, comprising:

conducting said plasma process in one of RIE mode, ICP mode, plasma mode and sputtering mode.

16. A method as recited in claim 1, comprising:

polishing respective first and second bonding surfaces to respective desired surface roughnesses and planarity; and

etching said first and second bonding surfaces after said polishing to activate said first and second bonding surfaces.

17. A method as recited in claim 16, wherein:

forming at least one of said first and second bonding surfaces comprises depositing a polishable material on a non-planar surface.

- 18. A method as recited in claim 17, wherein depositing said polishable material comprises depositing one of silicon oxide, silicon nitride or a dielectric polymer.
 - 19. A method as recited in claim 1, wherein said etching step comprises:

increasing available bonding energy of bonding pairs on said first and second bonding surfaces at approximately room temperature.

20. A method as recited in claim 19, comprising: obtaining a bond of at least 500 mJ/m².

- 21. A method as recited in claim 19, comprising: obtaining a bond of at least 1000 mJ/m².
- 22. A method as recited in claim 19, comprising: obtaining a bond of at least 2000 mJ/m².
- 23. A method as recited in claim 1, comprising: forming a bond of at least 500 mJ/m².
- 24. A method as recited in claim 1, comprising: obtaining a bond of at least 1000 mJ/m².
- 25. A method as recited in claim 1, comprising: obtaining a bond of at least 2000 mJ/m².
- 26. A method as recited in claim 1, comprising: forming chemical bonds between said bonding surfaces.
- 27. A method as recited in claim 26, comprising: forming chemical bonds between said bonding surfaces in one of ambient and vacuum.
- 28. A method as recited in claim 26, comprising: forming chemical bonds between said bonding surfaces in one of low and ultra high and vacuum.
- 29. A method as recited in claim 1, comprising: forming a bond of sufficient energy to virtually eliminate wafer bowing during subsequent processing of said bonded bonding surfaces.
- 30. A method as recited in claim 29, comprising:
 forming a bond of sufficient energy to virtually eliminate wafer bowing during subsequent thermal cycling of said bonded bonding surfaces.
- 31. A method as recited in claim 1, wherein said etching step comprises: increasing available bonding energy of bonding pairs on said first and second bonding surfaces at approximately room temperature; and

propagating said bonding at room temperature.

- 32. A method as recited in claim 31, comprising: propagating chemical bonding at room temperature.
- 33. A method as recited in claim 1, comprising:

depositing silicon dioxide as first and second bonding materials having said first and second bonding surfaces; and

etching said first and second bonding surfaces using an oxygen plasma.

34. A method as recited in claim 33, comprising:

rinsing said bonding materials in an ammonia-based solution after said etching.

35. A method as recited in claim 34, comprising:

rinsing said bonding materials in ammonium hydroxide after said etching.

36. A method as recited in claim 34, comprising:

rinsing said bonding materials in ammonium fluoride after said etching.

37. A method as recited in claim 1, comprising:

etching said first and second bonding surfaces under vacuum; and

bonding said first and second bonding surfaces without breaking said vacuum.

38. A method as recited in claim 1, comprising:

depositing a bonding material on each of first and second surfaces to obtain said first and second bonding surfaces.

39. A method as recited in claim 38, comprising:

depositing one of silicon dioxide and silicon nitride as said bonding material.

40. A method as recited in claim 1, comprising:

depositing silicon dioxide as said bonding material;

etching said silicon dioxide using one of oxygen, CF₄, and Ar plasma RIE; and

rinsing said silicon dioxide in an ammonia-based solution after said etching.

41. A method as recited in claim 1, comprising:

etching said first and second bonding materials using a wet etch process.

42. A method as recited in claim 42, comprising:

immersing said first and second bonding surfaces into a solution after said etching.

43. A method as recited in claim 1, comprising:

depositing silicon dioxide as said bonding material;

etching said silicon dioxide using one of diluted HF and diluted NH₄F.

44. A method as recited in claim 43, comprising:

rinsing said silicon dioxide in an ammonia-based solution after said etching.

45. A method as recited in claim 1, comprising:

forming said first and second bonding surfaces as silicon; and etching said bonding surfaces using a solution of HNO₃ and diluted HF.

46. A method as recited in claim 1, comprising:

forming said first and second bonding surfaces as silicon each having a native oxide layer; and

activating said native oxide layer using said etching step.

47. A method as recited in claim 1, comprising:

forming said first and second bonding surfaces each as silicon having a native oxide layer; and

exposing said first and second bonding surfaces to an oxygen plasma to etch said native oxide layers.

- 48. A method as recited in claim 47, creating a defective zone in said silicon during plasma etching.
- 49. A method as recited in claim 1, comprising: etching said first and second bonding materials using a plasma RIE process; forming a region having defects proximate to said bonding surface; and removing bonding by-products using said region.
- 50. A method as recited in claim 1, wherein said etching comprises: activating said first and second bonding surfaces; and creating a region under said first and second bonding surfaces for removing bonding by-products.
 - 51. A method as recited in claim 1, comprising:

creating a region proximate to said first and second bonding surfaces for at least one of removal and conversion of bonding by-products to a species capable of being absorbed by or diffusing away from said bonding surfaces.

52. A method as recited in claim 1, comprising:

forming said first bonding surface by depositing an oxide layer on a first semiconductor wafer;

forming said second bonding surface by depositing an oxide layer on a second semiconductor wafer; and

bonding said first and second semiconductor wafers.

53. A method as recited in claim 1, comprising:

forming said first bonding surface as a deposited oxide layer on a first semiconductor wafer, said first wafer comprising a first substrate and a first active region;

forming said second bonding surface as a deposited oxide layer on a second semiconductor wafer, said second wafer comprising a second substrate and a second active region;

bonding said first and second semiconductor wafers; and

removing at least a substantial portion of one of said first and second substrates after said bonding.

54. A method as recited in claim 1, comprising:

forming said first bonding surface as a deposited oxide layer on a semiconductor wafer, said wafer comprising a substrate and an active region;

forming said second bonding surface as a surrogate substrate;

bonding said wafer and said surrogate substrate; and

removing at least a substantial portion of said first substrate after said bonding.

- 55. A method as recited in claim 1, wherein said bonding comprises:
- maintaining contact between said first and second bonding surfaces for a specified period to produce bonding polymerization and allow removal of by-products.
 - 56. A method as recited in claim 55, comprising:

maintaining said contact between said first and second bonding surfaces for a period less than about 20 hours.

- 57. A method as recited in claim 1, wherein said bonding comprises:
- maintaining said first and second bonding surfaces for a specified period in ambient to remove bonding by-products.
 - 58. A method as recited in claim 1, comprising:

etching said first and second bonding surfaces using a bonding fixture under vacuum; bonding said first and second bonding surfaces using said fixture to bring together said first and second bonding surfaces while maintaining said vacuum.

59. A method as recited in claim 1, comprising:

forming a first oxide layer on a first wafer containing electrical devices; and polishing said first oxide layer to form said first bonding surface.

60. A method as recited in claim 59, comprising:

forming a second oxide layer on a second wafer containing electrical devices; and polishing said second oxide layer to form said second bonding surface.

61. A method as recited in claim 60, comprising:

forming said first oxide layer on said first wafer containing electrical devices of a first technology; and

forming said second oxide layer on said second wafer containing electrical devices of a second technology different from said first technology.

- 62. A method as recited in claim 60, comprising:
- interconnecting said first and second devices.
- 63. A method as recited in claim 60, comprising:

forming said first bonding surface on a surface of a one of a thermal spreader, surrogate substrate, antenna, wiring layer, and pre-formed multi-layer interconnect.

64. A method as recited in claim 60, comprising:

forming said second bonding surface on a surface of a one of a thermal spreader, surrogate substrate, antenna, wiring layer, and pre-formed multi-layer interconnect.

- 65. A method as recited in claim 1, comprising:
- forming said first bonding surface on a first wafer containing a first integrated circuit.
- 66. A method as recited in claim 65, comprising:

forming said second bonding surface on a second wafer containing a second integrated circuit.

67. A method as recited in claim 66, comprising:

forming said first bonding surface on said first wafer containing said first integrated circuit of a first technology; and

forming said second bonding surface on said second wafer containing said second integrated circuit of a second technology different from said first technology.

- 68. A method as recited in claim 66, comprising:
- interconnecting said first and second integrated circuits.
- 69. A method as recited in claim 1, comprising:
- said first and second bonding surfaces being substantially planar.
- 70. A method as recited in claim 1, wherein:

forming said first and second bonding surfaces comprises depositing a dielectric material.

71. A method as recited in claim 70, comprising:

polishing said dielectric material to a desired planarity and surface roughness.

72. A method as recited in claim 70, comprising:

depositing said dielectric material on a non-planar surface.

73. A method as recited in claim 72, wherein:

said polishing comprises chemical-mechanical polishing.

74. A method as recited in claim 1, comprising:

forming said first and second bonding surfaces to be non-planar; and polishing said first and second bonding surfaces to a desired planarity and surface roughness

75. A method as recited in claim 74, wherein:

said polishing comprises chemical-mechanical polishing.

76. A method as recited in claim 1, where said etching comprises:

activating said bonding surfaces; and

terminating said bonding surfaces with a desired species.

77. A method as recited in claim 1, wherein said etching comprises:

a first etching step to activate said bonding surfaces; and

a second etching step to terminate said bonding surfaces with a desired species.

78. A method as recited in claim 1, comprising:

obtaining etched bonding surfaces using said etching step; and

exposing said bonding surfaces to a gaseous chemical environment to terminate said etched bonding surfaces with a desired species.

79. A method as recited in claim 1, comprising:

forming a first oxide layer on a first wafer containing electrical devices and having a non-planar surface; and

forming a second oxide layer on a second wafer containing electrical devices; and polishing said first and second oxide layers to form said first and second bonding surfaces, respectively.

80. A method as recited in claim 79, comprising:

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forming said second oxide layer on said second wafer having a non-planar surface.

81. A method as recited in claim 1, comprising:

forming a first oxide layer on a first wafer containing electrical devices and having an irregular surface topology; and

forming a second oxide layer on a second wafer containing electrical devices; and polishing said first and second oxide layers to form said first and second bonding surfaces, respectively.

82. A method as recited in claim 81, comprising:

forming said second oxide layer on said second wafer having an irregular surface topology.

83. A bonding method, comprising:

forming first and second bonding surfaces each having a surface roughness in a range of 0.1 to 3 nm;

removing material from said first and second bonding surfaces while maintaining said surface roughness; and

directly bonding said first and second bonding surfaces at room temperature with a bonding strength of at least $500\ mJ/m^2$.

84. A method as recited in claim 83, comprising:

directly bonding said first and second bonding surfaces at room temperature with a bonding strength of at least 1000 mJ/m^2 .

85. A method as recited in claim 83, comprising:

directly bonding said first and second bonding surfaces at room temperature with a bonding strength of at least 2000 mJ/m^2 .

86. A method as recited in claim 83, comprising:

activating said first and second bonding surfaces and forming selected bonding groups on said first and second bonding surfaces.

87. A method as recited in claim 83, comprising:

polishing respective first and second bonding surfaces to said surface roughness; and etching said first and second bonding surfaces after said polishing to activate said first and second bonding surfaces.

88. A method as recited in claim 83, comprising:

converting bonding by-products to a species capable of being absorbed by or diffusing away from said bonding surfaces during said bonding step.

89. A method as recited in claim 83, comprising:

etching said first and second bonding surfaces using a plasma RIE process;

forming a subsurface layer having defects; and

removing bonding by-products using said subsurface layer.

90. A method as recited in claim 83, comprising:

forming said first bonding surface as a surface of a first semiconductor wafer having devices formed therein; and

forming said second bonding surface as a surface of a second semiconductor wafer having devices formed therein.

91. A method as recited in claim 90, comprising:

removing a substantial portion of said one of said first and second wafers.

92. A method as recited in claim 90, comprising:

interconnecting devices in said first and second wafers.

93. A method as recited in claim 83, comprising:

forming a first insulating layer on a first wafer containing electrical devices:

polishing said first insulating layer to form said first bonding surface;

forming a second insulating layer on a second wafer containing electrical devices; and polishing said second oxide layer to form said second bonding surface.

94. A method as recited in claim 83, comprising:

forming a first insulating layer on a first wafer containing electrical devices and having an irregular surface topology;

polishing said first insulating layer to form said first bonding surface;

forming a second insulating layer on a second wafer containing electrical devices and having an irregular surface topology; and

polishing said second oxide layer to form said second bonding surface.

95. A bonding method, comprising:

forming first and second bonding surfaces;

etching said first and second bonding surfaces;

terminating said first and second bonding surfaces with a species allowing formation of chemical bonds at about room temperature; and

bonding said first and second bonding surfaces at about room temperature.

96. A method as recited in claim 95, comprising:

bonding said first and second bonding surfaces at room temperature with a bonding strength of at least 500 mJ/m².

97. A method as recited in claim 95, comprising:

bonding said first and second bonding surfaces at room temperature with a bonding strength of at least 1000 mJ/m^2 .

98. A method as recited in claim 95, comprising:

bonding said first and second bonding surfaces at room temperature with a bonding strength of at least 2000 mJ/m^2 .

99. A method as recited in claim 95, comprising:

activating said first and second bonding surfaces prior to said bonding step.

100. A method as recited in claim 95, comprising:

polishing said first and second bonding surfaces; and

etching said first and second bonding surfaces after said polishing to activate said first and second bonding surfaces.

101. A method as recited in claim 95, comprising:

converting bonding by-products to a species capable of being absorbed by or diffusing away from said bonding surfaces during said bonding step.

102. A method as recited in claim 95, comprising:

forming said first bonding surface as a surface of a first semiconductor wafer having devices formed therein; and

forming said second bonding surface as a surface of a second semiconductor wafer having devices formed therein.

103. A method as recited in claim 102, wherein one of said first and second wafers comprises a substrate, said method comprising:

removing a substantial portion of said one of said first and second wafers.

104. A method as recited in claim 102, comprising:

interconnecting devices in said first and second wafers.

105. A method as recited in claim 95, comprising:

forming a first insulating layer on a first wafer containing electrical devices; polishing said first insulating layer to form said first bonding surface; forming a second insulating layer on a second wafer containing electrical devices; and polishing said second oxide layer to form said second bonding surface.

106. A method as recited in claim 95, comprising:

forming a first insulating layer on a first wafer containing electrical devices and having an irregular surface topology;

polishing said first insulating layer to form said first bonding surface;

forming a second insulating layer on a second wafer containing electrical devices and having an irregular surface topology; and

polishing said second oxide layer to form said second bonding surface.

107. A bonded device, comprising:

a first material having a first etched bonding surface; and

a second material having a second etched bonding surface directly bonded to said first bonding surface at room temperature having a bonding strength of at least 500 to 2000 mJ/m².

108. A device as recited in claim 107, comprising:

said first and second bonding surfaces being activated and terminated with a desired bonding species.

- 109. A device as recited in claim 108, wherein said desired species comprise:
- a monolayer of one of a desired atom and a desired molecule on said bonding surface.
- 110. A device as recited in claim 108, wherein said desired species comprise at least one of a silanol group, an NH₂ group, a fluorine group and an HF group.
 - 111. A device as recited in claim 107, comprising:

said first and second bonding surfaces each having a defective region located proximate to said first and second bonding surfaces, respectively.

112. A device as recited in claim 107, wherein:

said first material comprises a surface of a first semiconductor wafer having devices formed therein; and

said second material comprises a surface of a second semiconductor wafer having devices formed therein.

- 113. A device as recited in claim 112, wherein one of said first and second wafers comprises a device region after removing a substantial portion of a substrate of said one of said first and second wafers.
 - 114. A device as recited in claim 112, comprising:

devices in said first and second wafers being interconnected.

115. A device as recited in claim 112, comprising:

said first and second wafers being different technologies.

116. A device as recited in claim 107, wherein:

one of said first and second wafers comprises an integrated circuit.

117. A device as recited in claim 116, comprising:

devices in said first and second wafers being interconnected.

118. A device as recited in claim 116, comprising:

said first and second wafers having an irregular surface topology.

119. A device as recited in claim 107, wherein:

said first material comprises a first wafer containing electrical devices and having a first non-planar surface; and

said first bonding surface comprises a polished and etched deposited oxide layer on said first non-planar surface.

120. A device as recited in claim 119, wherein:

said second material comprises a second wafer containing electrical devices and having a second non-planar surface; and

said second bonding surface comprises a polished, planarized and etched deposited oxide layer on said second non-planar surface.

121. A method as recited in claim 107, wherein:

said first material comprises a first wafer containing electrical devices and having a first surface with irregular topology; and

said first bonding surface comprises a polished, planarized and etched deposited oxide layer on said first surface.

122. A method as recited in claim 121, comprising:

said second material comprises a second wafer containing electrical devices and having a second surface with irregular topology; and

said second bonding surface comprises a polished, planarized and etched deposited oxide layer on said second surface.

- 123. A bonded device, comprising:
- a first material having a first etched and activated bonding surface terminated with a first desired bonding species; and
- a second material having a second etched and activated bonding surface terminated with a second desired bonding species bonded to said first bonding surface at room temperature.
- 124. A device as recited in claim 123, wherein said species comprise at least one of a silanol group, an NH₂ group, a fluorine group and an HF group.
 - 125. A device as recited in claim 123, comprising:
- said first and second bonding surfaces each having a defective region located proximate to said surfaces.
 - 126. A device as recited in claim 123, wherein said desired species comprises: a monolayer of one of a desired atom and a desired molecule on said bonding surface.
 - 127. A device as recited in claim 123, comprising:
- said second bonding surface bonded to said first bonding surface at room temperature having a bonding strength of at least 500 to 2000 mJ/m².
 - 128. A device as recited in claim 123, wherein:
- said first bonding surface comprises a surface of a first semiconductor wafer having devices formed therein; and
- said second bonding surface comprises a surface of a second semiconductor wafer having devices formed therein.
- 129. A device as recited in claim 128, wherein one of said first and second wafers comprises a device region after removing a substantial portion of a substrate of said one of said first and second wafers.
 - 130. A device as recited in claim 123, comprising: devices in said first and second wafers being interconnected.
 - 131. A device as recited in claim 123, comprising:

said first and second wafers being different technologies.

132. A device as recited in claim 123, wherein:

one of said first and second wafers comprises an integrated circuit.

133. A device as recited in claim 123, comprising:

devices in said first and second wafers being interconnected.

134. A device as recited in claim 123, wherein:

said first material comprises a first wafer containing electrical devices and having a first non-planar surface; and

said first bonding surface comprises a polished and etched deposited oxide layer on said first non-planar surface.

135. A device as recited in claim 134, wherein:

said second material comprises a second wafer containing electrical devices and having a second non-planar surface; and

said second bonding surface comprises a polished, planarized and etched deposited oxide layer on said second non-planar surface.

136. A method as recited in claim 123, wherein:

said first material comprises a first wafer containing electrical devices and having a first surface with irregular topology; and

said first bonding surface comprises a polished, planarized and etched deposited oxide layer on said first surface.

137. A method as recited in claim 123. wherein:

said second material comprises a second wafer containing electrical devices and having a second surface with irregular topology; and

said second bonding surface comprises a polished, planarized and etched deposited oxide layer on said second surface.

138. A method as recited in claim 1, wherein said first and second bonding surfaces are respective surfaces of first and second substrates, the method comprising:

removing a substantial portion of one of said first and second substrates.

139. A method as recited in claim 138, comprising:

annealing said bonded substrates after said removing step.

140. A method as recited in claim 138, wherein said first and second substrates have

first and second electrical devices, the method comprising:

interconnecting said first and second devices after said removing step.

141. A method as recited in claim 91, comprising:

annealing said bonded substrates after said removing step.

142. A method as recited in claim 91, comprising:

interconnecting devices in said first and second semiconductor wafers after said removing step.

143. A method as recited in claim 103, comprising:

annealing said bonded substrates after said removing step.

144. A device as recited in claim 113, wherein:

said device region comprises an annealed device region.

145. A device as recited in claim 129, wherein;

said first and second materials comprise respective first and second substrates; and one of said first and second substrates comprises a substrate region obtained by removing a substantial portion of a substrate of said one of said first and second substrate.

146. A device as recited in claim 145, wherein:

said substrate region comprises an annealed substrate region.

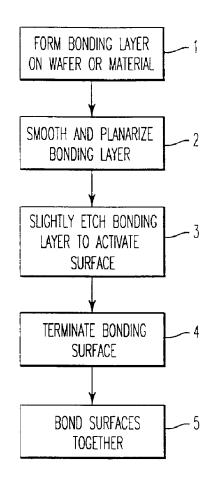


FIG. 1

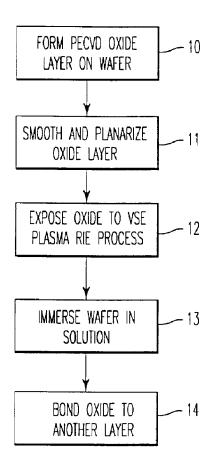
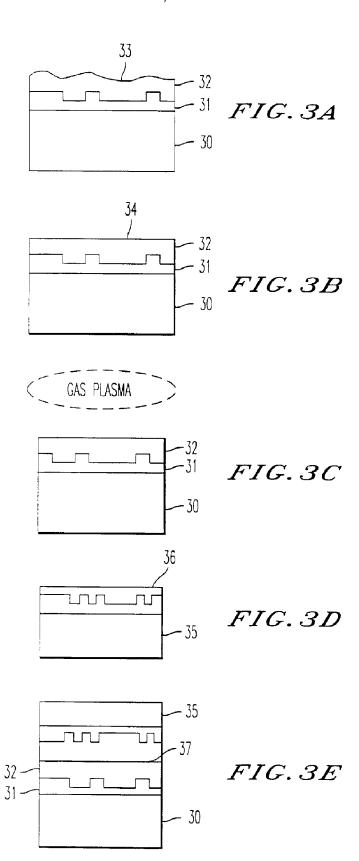
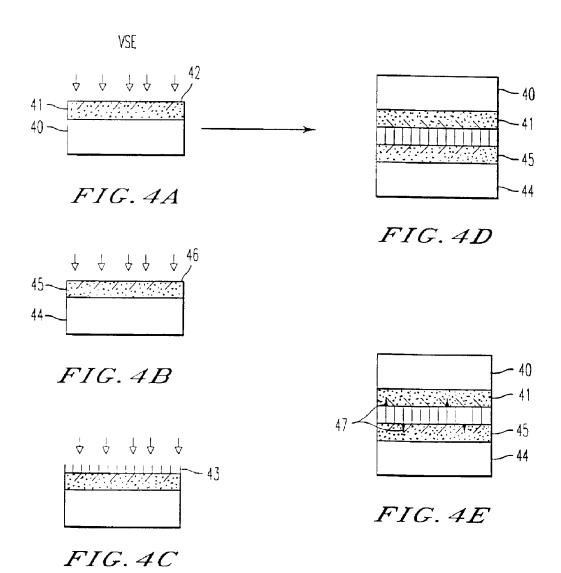
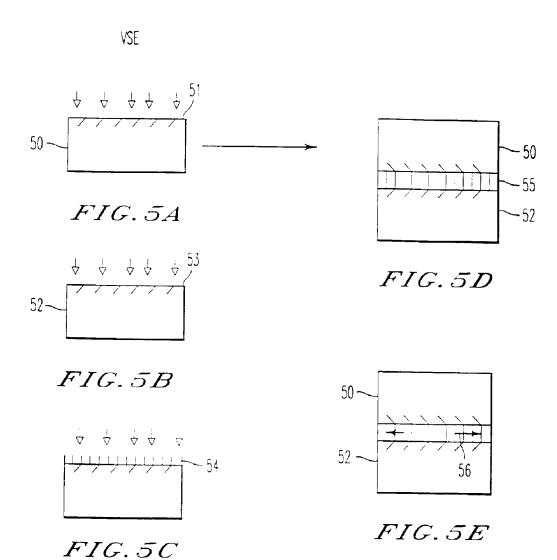


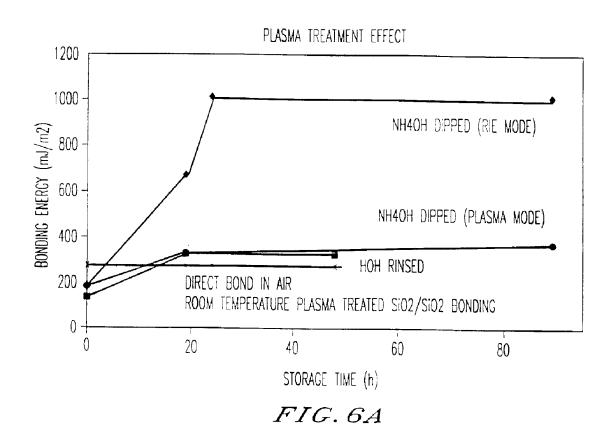
FIG. 2



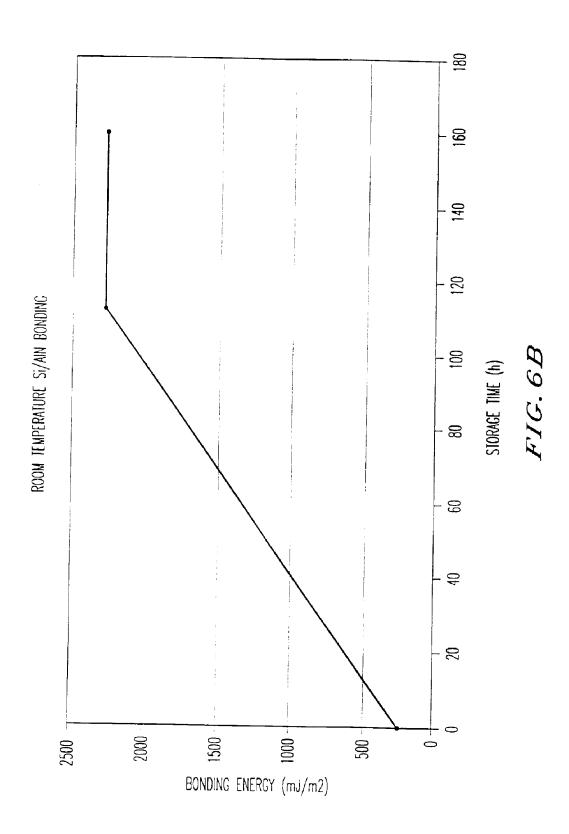
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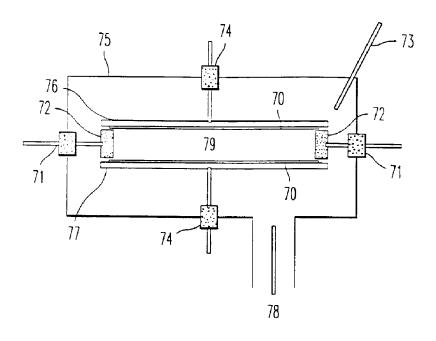
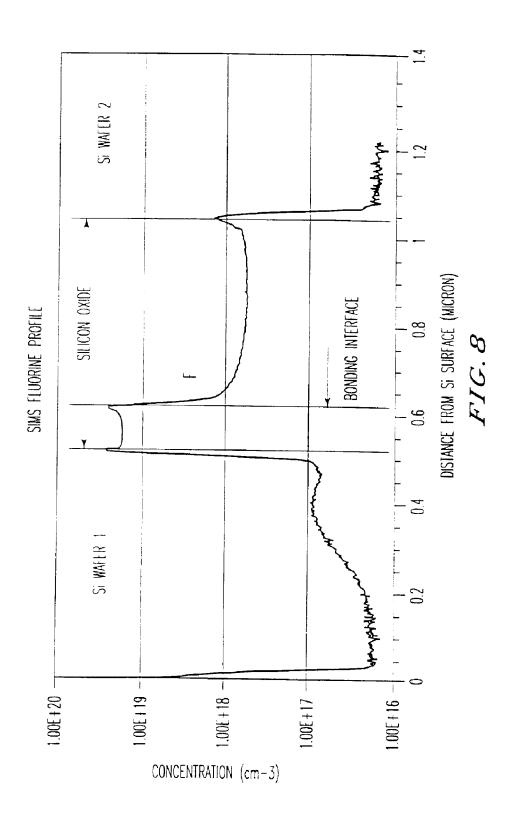


FIG. 7



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(19)日本国特許庁(JP)

(12) 公開特許公報(A)

(11)特許出願公開番号

特開平8-195334

(43)公開日 平成8年(1996)7月30日

(51) Int.Cl. ⁶		識別記号	庁内整理番号	FΙ	技術表示箇所
H01L	21/02	В			
	21/308	G			
	27/12	В			

審査請求 未請求 請求項の数8 OL (全 5 頁)

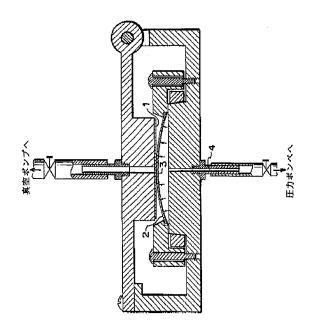
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(54) 【発明の名称】 シリコン基板の接合方法

(57)【要約】

【目的】 シリコン基板どうしの接合工程において、低温処理で強固な接合を可能とすることにあり、また、これにより高温に耐える材料以外の材料も利用可能とし、更に膜はがれや、ボイドの発生も防止する。

【構成】 シリコン基板 1, 2表面を疎水化処理する工程と、該疎水化処理した前記基板どうしを加圧下で接触させて接合する工程と、その後加熱処理して前記接合を強化する工程と、を有することを特徴とするシリコン基板の接合方法。前記基板どうし1, 2を加圧下で接触させて接合する工程は、前記基板 1, 2の中央部から順次接触させることを特徴とする。



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【特許請求の範囲】

【請求項1】 シリコン基板表面を疎水化処理する工程と、該疎水化処理した前記基板どうしを加圧下で接触させて接合する工程と、その後加熱処理して前記接合を強化する工程と、を有することを特徴とするシリコン基板の接合方法。

【請求項2】 前記疎水化処理は、フッ酸により前記基板表面の自然酸化膜を除去することを特徴とする請求項1に記載のシリコン基板の接合方法。

【請求項3】 前記疎水化処理は、バッファーフッ酸を 用いることを特徴とする請求項1に記載のシリコン基板 の接合方法。

【請求項4】 前記疎水化処理は、高真空中で基板表面に加速した不活性イオンを照射し、清浄なシリコン表面を形成した後、水素ガスを導入して基板表面を水素原子で終端することを特徴とする請求項1に記載のシリコン基板の接合方法。

【請求項5】 前記基板どうしを加圧下で接触させて接合する工程は、前記基板の中央部から順次接触させることを特徴とする請求項1に記載のシリコン基板の接合方法。

【請求項6】 前記加熱処理は250℃~700℃で行なうことを特徴とする請求項1に記載のシリコン基板の接合方法。

【請求項7】 前記加熱処理は、前記シリコン基板の中央より温度上昇させることを特徴とする請求項1に記載のシリコン基板の接合方法。

【請求項8】 前記加熱処理は、前記シリコン基板の端部より温度上昇させることを特徴とする請求項1に記載のシリコン基板の接合方法。

【発明の詳細な説明】

[0001]

【産業上の利用分野】本発明は、半導体基板の製造方法 に関し、特にシリコン基板どうしの接合方法に関する。

[0002]

【従来の方法】シリコン基板どうしの接合は、SOI(Silicon on Insulator)基板の作製に不可欠な技術である。従来、シリコン基板同士の接合は、特開昭39-17869号公報あるいは特開昭62-27040号公報に見られるように、基板を接触後、酸化雰囲気中で加熱する、あるいは親水化処理した基板を接触後加熱する方法が用いられており、基板は酸化層を介して接合されている。

[0003]

【発明が解決しようとする課題】しかしながら、これまでの方法では、強固な接合を得るためには、接触後 1 0 0 0 ℃前後に加熱する必要がある。従って、接合前に基板上に素子が形成されているような場合、このような高温に耐えるためには使用する材料が極めて制限されるという問題がある。

【0004】さらに、積層された構造においては各層材料の熱膨張差により膜剥がれが発生するという問題もある。従来の方法はまた、基板表面の水酸基の脱水分解により発生したガスがボイドとして残りやすく、これが接合強度を低下させるという問題もあった。

【0005】(発明の目的)本発明の目的は、シリコン基板の接合工程において、低温処理で強固な接合を可能とすることにあり、また、これにより高温に耐える材料以外の材料も利用可能とし、更に膜はがれや、ボイドの発生も防止することにある。

[0006]

【課題を解決するための手段】本発明のシリコン基板の接合方法は、前記課題を解決するための手段としてシリコン基板表面の疎水化処理工程、加圧しながらの基板どうしの接触工程、加熱処理工程の3つの工程を有することを特徴とする。

【0007】まず、シリコン表面の疎水化は、代表的にはフッ酸による表面の自然酸化膜を除去することでおこなわれる。フッ酸処理によりシリコン表面の大部分が水素原子で終端される。使用されるフッ酸の濃度は10重量%以下であり、1~5%のフッ酸処理が疎水化処理に適している。PHを3~6に調整したバッファーフッ酸を使用してもよい。またフッ酸処理前に基板表面に付着している有機物等の不純物を除去する処理をしてもよい。この場合、基板表面に凹凸を形成するような処理は避けなければならない。疎水化処理方法としては上記フッ酸処理の他、高真空中で基板表面に加速した不活性イオンを照射し、清浄なシリコン表面を形成した後、水素ガスを真空槽に導入し、基板表面を水素原子で終端する方法がある。

【0008】次に、疎水化処理された2枚の基板は圧力を加えながら接触させられる。図1は本発明で基板の接触に使用した装置の概略図である。この装置は第1の基板1を平坦に保持し、流体導入口4から加圧された流体(液体、気体のどちらでもよい)を導入し、この流体圧を第2の基板2の裏面に印加しながら2枚の基板を近接させるものである。図1の装置では流体圧は薄いメンブレン3に加えられ、このメンブレンが球面状にたわむため、それに接する第2基板2も球面状にたわむ。第2基板2がたわむことにより2枚の基板はそれぞれの中央部から接触していく。この接触方法では、接触過程で雰囲気ガスが周辺部に追い出されるため、大気中においても基板間にガスが残留することがない。もちろん2枚の基板を真空中で接触させてもよい。流体圧は3kg/cmプ以上あれば十分効果がある。

【0009】次に、接触した2枚のシリコン基板の接合を強固にするために加熱する。本発明では加熱温度が250℃以上あれば実用上十分な接合強度が得られるが、300℃~700℃、さらに好ましくは500~700℃であれば従来方法に匹敵する強固な接合が達成され

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る。ヒータ加熱、ランプ加熱等の加熱手段により接合基板を昇温する。また、加熱中の雰囲気は接合には制限がなく大気中、不活性ガス中、真空中いずれでもよい。

【0010】加熱する効果は以下のように推測される。 加熱によりシリコン基板表面に吸着している水素原子が 脱離することにより、基板最表面にあるシリコン原子同 士が直接結合するか、あるいは水素原子と大気中の酸素 原子が置き変わり、酸素原子を介してシリコン原子が結 合することによって2枚のシリコン基板の接合がおこな われる。

【0011】加熱中における接合強度の増加過程において、基板周辺部より強い結合がすすんだ場合において、基板の中央部にしばしば空孔(ボイド)が発生することがある。一旦強固に接合した部分はひずみを緩和することができず、未接合部にひずみが集中した結果ボイドとして残るものと考えられる。こうしたボイドは基板温度の不均一性により強固に接合した部分がランダムに作られたときにも発生する。

【0012】したがって、ボイドを発生させずにシリコン基板を接合させるためには一点より接合をすることが有効である。その方法として、図3に示すように接触が完了したシリコン基板5を石英やセラミクス等熱伝導率が比較的小さい2枚の板6および6′で挟み、これらの板の背後の基板中央を加熱することが有効である。こうした手段により2枚のシリコン基板は中央より徐々に加熱され、強固な接合部が中央より周辺に広がりボイドの発生が抑えられる。

【0013】他の方法として図4に示すように、接触させたシリコン基板8を搬送ローラ10によって細長い加熱手段9、9′の短辺方向に搬送させ、基板の端より徐30々に強固な接合部を形成する方法がある。遮蔽板12および冷却ローラ11は未接合部への熱伝導を低下させることにより温度上昇部を制御する働きをする。図3、図4いずれの方法においても加熱手段を上下2組設置することにより、2枚の基板の温度差をできるだけ小さくしている。また、図4に示した方法においては基板を搬送するかわりに加熱手段を基板の端より移動させても同様の効果が期待できる。

[0014]

【作用】本発明によれば、あらかじめ基板表面を疎水化 40 処理しておくため、低温度でも強固な接合が可能となる。

【0015】また、前記疎水化処理は、フッ酸、バッファーフッ酸により行なうことにより、疎水化処理を効果的に行なうことができる。

【0016】また、前記疎水化処理は、高真空中で基板表面に加速した不活性イオンを照射し、清浄なシリコン表面を形成した後、水素ガスを導入して基板表面を水素原子で終端することにより、基板表面の疎水化を完全にすることができる。

【0017】また、前記基板どうしを加圧下で接触させて接合する工程は、前記基板の中央部から順次接触させることにより、接合界面に基体等を含むことがなくなり、ボイドの発生を防止できる。

【0018】また、前記加熱処理は250℃ \sim 700℃ で行なうことにより、十分な接合強度を得ることができる。

[0019]

【実施例】

10 [実施例1] 2枚の(100)シリコン基板を2%フッ酸で表面疎水化処理し、ただちに図1の接触装置により5気圧のガス圧を印加しつつ、大気中で2枚のシリコン基板を接触させた。

【0020】つぎに大気中で2時間加熱して接合した。 加熱温度(熱処理温度)は、図2に示した各温度とした。

【0021】こうして作製した接合試料の曲げ強度を測定した。図2は曲げ強度より算出した破壊時の界面せん断応力を図示したものである。同様の方法で測定したシリコン基板の破壊強度は55 k g/c m² であり、25 0 $^{\circ}$ で加熱すれば十分強固に接合されていることがわかった。

【0022】 [実施例2] (100) シリコン基板を200℃で10分間紫外線処理し、基板表面の有機不純物を除去した後、フッ化アンモニウムでPH4に調整したバッファーフッ酸で疎水化処理した。

【0023】その後実施例1と同様に2枚の基板を接触させた後、加熱処理を行った。

【0024】実施例1と同様にして強度を測定したところ、同等の強度で接合することが確かめられた。

【0025】 [比較例] (100) シリコン基板を親水 化処理し、実施例1と同様に接触、加熱処理をした試料 の接合強度を測定したところ、図2に示したように、6 00℃以上では強固に接合したが、低温では十分な強度 が得られなかった。また、400℃以下の場合、ダイシ ングソー切断時において接合面で剥離した。

【0026】 [実施例3] 2枚の(100)シリコン基板を85℃の硫酸および過酸化水素混合水溶液中に5分浸漬した後、10分間純水により洗浄した。ひきつづき、2%フッ酸水溶液で20秒間疎水化処理を行い、2分間純水洗浄した。以上の前処理を施した2枚のシリコン基板を図1に示した接触装置により5気圧を印加しつつ接触させた。次に図3に概略を示すように、接触した2枚のシリコン基板5を2枚のアルミナセラミクス板6、6′に挟み、このアルミナセラミクス板の中央部を加熱温度が500℃となるようにヒータ7、7′により加熱した。

【0027】以上のように接合した2枚のシリコン基板に赤外線を照射し、透過光を赤外線カメラにより観察したところ、ボイドの発生は見られなかった。また、破壊

5

時のせん断応力は $85kg/cm^2$ であり十分な強度をもつことが確認された。

【0028】 [実施例4] 2枚の(100) シリコン基板 を酸素雰囲気中で200℃に加熱しながら、紫外線を2 0分間照射したのち、2%フッ酸水溶液で20秒間疎水 化処理を行い、2分間純水洗浄した。以上の疎水化処理 を施した2枚のシリコン基板を図1に示した接触装置に より5気圧を印加しつつ接触させた。次に図4に概略を 示す試料搬送装置を有する加熱装置により接触したシリ コン基板8を加熱した。図4において、10は搬送用ロ ーラ、11は水冷されたローラであり、12は熱遮蔽板 である。シリコン基板8は矢印方向に移動する過程で熱 遮蔽版12の中央部に設置されたヒータ9、9′により 加熱される。シリコン基板のヒータ直下部分は温度上昇 するが、そこからはなれた部分は水冷ローラにより熱が 遮断され低温に保たれるため、強固な接合は起こらな い。本実施例においてはヒータ直下でのシリコン基板温 度が550℃となるようにヒータ電力を調整した。

【0029】以上のように接合したシリコン基板を実施例3と同様に赤外線カメラで観察したがボイドの発生は20見られなかった。また、破壊時のせん断応力は91kg/ cm^2 であった。

[0030]

【発明の効果】以上説明したように、本発明の接合方法は、従来に比べ低温でシリコン基板の強固な接合を可能*

*にする効果が得られる。また、本発明の方法によれば、 低温で接合可能であるため、素子形成後のシリコン基板 どうしも接合することができ、マイクロマシン等の立体 構造体の形成に有効な手段を提供することができる。

【0031】また、積層構造の熱膨張差による膜はがれや、ボイドの発生等も、少なくすることができるため、 品質や、信頼性も向上できる。

【図面の簡単な説明】

【図1】本発明で基板の接触に使用した装置の概略図

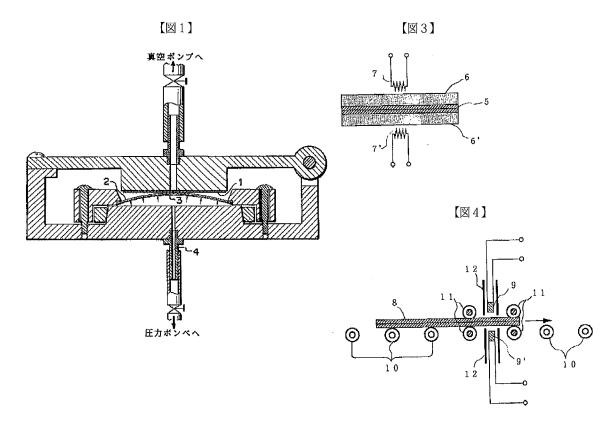
【図2】曲げ強度より算出した実施例と比較例の破壊時の界面せん断応力の図

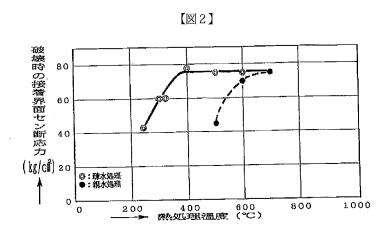
【図3】実施例3で用いた加熱方法の概略図

【図4】実施例4で用いた加熱装置の概略図

【符号の説明】

- 1 第1の基板
- 2 第2の基板
- 3 メンブレン
- 4 流体導入口
- 5、8 接触させたシリコン基板
- 6、6′、セラミクス板
- 7、7′、9、9′ 加熱手段
- 10 搬送ローラ
- 11 水冷ローラ
- 12 熱遮蔽板





(11)Publication number

08-195334

(43)Date of publication of application

30.07.1996

(51)Int.Cl.

H01L 21/02

H01L 21/308

H01L 27/12

(21)Application number

07-040398

(22)Date of filing

28.02.1995

(71)Applicant

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(30) Priority

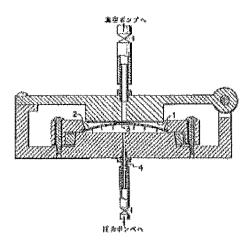
Priority number: 06283470 Priority date: 17.11.1994 Priority country: JP

(54)METHOD FOR BONDING SILICON SUBSTRATE

(57) Abstract

PURPOSE: To firmly bond silicon substrates at low temperature, to utilize materials which cannot stand high temperature and to prevent films from separating and voids from being produced.

CONSTITUTION: The method comprises the steps of hydrophobing the surfaces of silicon substrates 1 and 2, contacting both hydrophobed silicon substrates, press-bonding them, and heating them to strengthen the bond. And the press-bonding step, the substrates 1 and 2 are brought into contact from the central portions of the substrates 1 and 2 to the peripheries.



CLAIMS

[Claim(s)]

[Claim 1]A joining method of a silicon substrate characterized by comprising the following.

A process of carrying out hydrophobing processing of the silicon substrate surface.

A process which the aforementioned substrates which carried out hydrophobing processing are contacted under application of pressure, and is joined, and its process of carrying out afterbaking processing and strengthening the aforementioned junction.

[Claim 2]A joining method of the silicon substrate according to claim 1, wherein the aforementioned hydrophobing processing removes a natural oxide film of the aforementioned substrate surface by fluoric acid.

[Claim 3]A joining method of the silicon substrate according to claim 1, wherein the aforementioned hydrophobing processing uses buffer fluoric acid.

[Claim 4]A joining method of the silicon substrate according to claim 1 after irradiating inactive ions accelerated to a substrate surface in a high vacuum and forming a pure silicon surface, wherein the aforementioned hydrophobing processing introduces hydrogen gas and carries out the termination of the substrate surface with a hydrogen atom.

[Claim 5]A joining method of the silicon substrate according to claim 1 contacting sequentially a process which the aforementioned substrates are contacted under application of pressure, and is joined from a central part of the aforementioned substrate.

[Claim 6]A joining method of the silicon substrate according to claim 1 performing the aforementioned heat-treatment at 250 degrees C - 700 degrees C.

[Claim 7]A joining method of the silicon substrate according to claim 1 carrying out temperature rise of the aforementioned heat-treatment from a center of the aforementioned silicon substrate.

[Claim 8]A joining method of the silicon substrate according to claim 1 carrying out temperature rise of the aforementioned heat-treatment from an end of the aforementioned silicon substrate.

DETAILED DESCRIPTION

[Detailed Description of the Invention]

[0001]

[Industrial Application] The present invention relates to the manufacturing method of a semiconductor substrate, and relates especially to the joining method of silicon substrates.

[0002]

[The conventional method] Junction of silicon substrates is technology indispensable to production of a SOI (Silicon on Insulator) substrate. A method of carrying out contact afterbaking of the substrate which heated or carried out hydrophilization treatment of the substrate in an oxidizing atmosphere after contacting is used so that junction of silicon substrates may be conventionally looked at by JP,S39-17869,A or JP,S62-27040,A.

A substrate is joined via an oxidizing zone.

[0003]

[Problem to be solved by the invention] However, in an old method, in order to acquire firm junction, it is necessary to heat at around 1000 degrees C after contact. Therefore, when the element is formed on the substrate before junction, in order to bear such an elevated temperature, there is a problem that the material to be used is restricted extremely.

[0004] There is also a problem that film peeling occurs according to the thermal expansion difference of class material in the laminated structure. The gas emitted by drying disassembly of the hydroxyl group of a substrate surface remained easily as a void again, and the conventional method also had the problem of this making bonding strength be deteriorated.

[0005] (The object of invention) There is the object of this invention in supposing that materials other than the material which is in enabling firm junction by low temperature treatment, and bears an elevated temperature by this are also available, and also preventing film peeling and generating of a void in the joining process of a silicon substrate.

[0006]

[Means for solving problem] The joining method of the silicon substrate of the present invention is characterized by hydrophobing down stream processing of a silicon substrate surface, and having three processes of the contact process of substrates while pressurizing, and a heat-treatment process as above-mentioned The means for solving a technical problem.

[0007] First, hydrophobing of a silicon surface is typically performed by removing the natural oxide film of the surface by fluoric acid. The termination of most silicon surfaces is carried out by fluoric acid processing with a hydrogen atom. The concentration of the fluoric acid used is 10 % by weight or less, and 1 to 5% of fluoric acid processing is suitable for hydrophobing processing. The buffer fluoric acid which adjusted PH to 3·6 may be used. Processing which removes impurities, such as an organic matter which has adhered to the substrate surface before fluoric acid processing, may be carried out. In this case, processing which forms unevenness in a substrate surface must be avoided. After irradiating the inactive ions accelerated to the substrate surface in the high vacuum besides the above-mentioned fluoric acid processing as a hydrophobing processing method and forming a pure silicon surface, hydrogen gas is introduced into a vacuum chamber and there is the method of carrying out the termination of the substrate surface with a hydrogen atom.

[0008]Next, the substrate of two sheets by which hydrophobing processing was carried out is contacted while applying a pressure. Fig.1 is a schematic view of the equipment used for contact of a substrate by the present invention. This equipment holds the first substrate 1 evenly, introduces the fluid (either a liquid or a gas is OK) pressurized from the fluid introducing port 4, and it makes the substrate of two sheets approach, applying this hydrostatic pressure to the back surface of the second substrate 2. Since hydrostatic pressure is applied to the thin membrane 3 and this membrane is bent by the equipment of Fig.1 in sphere form, the 2nd substrate 2 that touches it is also bent by it in sphere form. When the 2nd substrate 2 bends, the substrate of two sheets contacts from each central part. Since a controlled atmosphere is driven out by the periphery in a contact process in this contact method, into the atmosphere, gas does not have a residual thing [carrying out] between substrates. Of course, the substrate of two sheets may be contacted in a vacuum. If, as for hydrostatic pressure, there is more than 3 kg/cm², it is effective enough.

[0009]Next, it heats in order to strengthen junction of the silicon substrate of two sheets which contacted. In the present invention, if the heating temperature is 250 degrees C or more, practically sufficient bonding strength will be obtained, but 300 degrees C - 700 degrees C, if it is 500-700 degrees C still more preferably, the firm junction which is equal to the conventional method will be attained. The temperature rise of the joined substrate is carried out by heating means, such as heater heating and lamp heating. Restriction may not be in junction and any may be sufficient as the atmosphere under heating among the atmosphere, inactive gas, and a vacuum.

[0010] The effect to heat is guessed as follows. When the hydrogen atom which is sticking to a silicon substrate surface by heating disconnects itself, The silicon atoms in the substrate outermost surface couple directly, or a hydrogen atom and the oxygen atom in the atmosphere place and change, and when a silicon atom joins together via an oxygen atom, junction of the silicon substrate of two sheets is performed.

[0011]In the increasing process of the bonding strength under heating, when a strong combination progresses from a substrate periphery, a hole (void) may often occur in the central part of the substrate. It is thought that the portion once joined firmly remains as a void as a result of being unable to ease a strain but a strain concentrating on an unsealed part. Such a void occurs, also when the portion firmly joined by the heterogeneity of substrate temperature is made at random.

[0012] Therefore, in order to join a silicon substrate, without generating a void, joining from one point is effective. It is effective to pinch the silicon substrate 5 which contact completed as the method, by the board 6 and 6' which are two sheets whose thermal conductivity, such as quartz and ceramics, is comparatively small, as shown in Fig.3, and to heat the center of a substrate of these boards in back. The silicon substrate of two sheets is gradually heated from a center by such a means, a firm junction spreads on a periphery from a center, and generating of a void is suppressed.

[0013]As shown in Fig.4 as other methods, the short side direction of the long and slender heating means 9 and 9' is made to convey the contacted silicon substrate 8 with the conveying roller 10, and there is the method of forming a firm junction gradually from the end of a substrate. The shield 12 and the cooling roller 11 serve to control a temperature rise part by making heat conduction to an unsealed part be deteriorated. The temperature gradient of the substrate of two sheets is made as small as possible by 2 sets of upper and lower sides installing a heating means also in which method of Fig.3 and Fig.4. The same effect is expectable even if it moves a heating means from the end of a

substrate instead of conveying a substrate in the method shown in Fig.4.

[0014]

[Function] According to the present invention, in order to carry out hydrophobing processing of the substrate surface beforehand, firm joining becomes possible also for low temperature.

[0015] Hydrophobing processing can be effectively performed by performing the aforementioned hydrophobing processing by fluoric acid and buffer fluoric acid.

[0016] After the aforementioned hydrophobing processing irradiates the inactive ions accelerated to the substrate surface in the high vacuum and forms a pure silicon surface, it can carry out hydrophobing of a substrate surface completely by introducing hydrogen gas and carrying out the termination of the substrate surface with a hydrogen atom.

[0017] By making it contact sequentially from the central part of the aforementioned substrate, it is lost that a base substance etc. are included of the process which the aforementioned substrates are contacted under application of pressure, and is joined in a bonding interface, and it can prevent generating of a void.

[0018]Sufficient bonding strength can be obtained by performing the aforementioned heat-treatment at 250 degrees C - 700 degrees C.

[0019]

[Working example]

[Working example 1] The silicon substrate of two sheets was contacted in the atmosphere, carrying out surface hydrophobing processing of (100) the silicon substrate of two sheets by fluoric acid 2%, and applying 5-atmosphere gas pressure with the contactor of Fig.1 immediately.

[0020]Next, in the atmosphere, it heated for 2 hours and joined. Heating temperature (heat treatment temperature) was made into each temperature shown in Fig.2.

[0021]In this way, the flexural strength of the produced junction sample was measured. Fig.2 illustrates the interface shearing stress at the time of the destruction computed from flexural strength. The disruptive strength of the silicon substrate measured in the similar way was 55 kg/cm², and when heating at 250 degrees C, it turned out that it is joined sufficiently firmly.

[0022] [Working example 2] (100) After carrying out ultraviolet treatment of the silicon substrate for 10 minutes at 200 degrees C and removing the organic impurities of a substrate surface, hydrophobing processing was carried out by the buffer fluoric acid adjusted to PH4 with ammonium fluoride.

[0023] Heat-treatment was performed after contacting the substrate of two sheets like the working example 1 after that.

[0024] When strength was measured like the working example 1, joining by equivalent strength was confirmed.

[0025][Comparative example] (100) Hydrophilization treatment of the silicon substrate was carried out, when the bonding strength of the sample which carried out contact and heat-treatment like the working example 1 was measured, as shown in Fig.2, above 600 degrees C, it joined firmly, but strength sufficient at low temperature was not obtained. In the case of 400 degrees C or less, it exfoliated in the plane of composition at the time of dicing saw cutting.

[0026][Working example 3] After (100) the silicon substrate of two sheets was immersed for 5 minutes into sulfuric acid and a hydrogen peroxide mixed aqueous solution of 85 degrees C, pure

water washed for 10 minutes. It pulled and continued, and performed and carried out pure water washing of the hydrophobing processing for 2 minutes for 20 seconds in the 2% fluoric acid aqueous solution. It was made to contact, applying 5 atmospheres with the contactor which showed the silicon substrate of two sheets which pretreated more than to Fig.1. Next, as an outline was shown in Fig.3, the two alumina ceramic boards 6 and 6' pinched the silicon substrate 5 of two sheets which contacted, and the central part of this alumina ceramic board was heated by the heater 7 and 7' so that heating temperature might be 500 degrees C.

[0027] Generating of the void was not seen, when infrared rays were irradiated to the silicon substrate of two sheets joined as mentioned above and the transmitted light was observed with the infrared camera. The shearing stress at the time of destruction is 85 kg/cm², and having sufficient strength was checked.

[0028] [Working example 4] After irradiating ultraviolet rays for 20 minutes, heating (100) the silicon substrate of two sheets at 200 degrees C in oxygen environment, pure water washing of the hydrophobing processing was performed and carried out for 2 minutes for 20 seconds in the 2% fluoric acid aqueous solution. It was made to contact, applying 5 atmospheres with the contactor which showed the silicon substrate of two sheets which performed the above hydrophobing processing to Fig. 1. Next, the silicon substrate 8 which contacted with the heating apparatus which has a sample conveying machine which shows an outline to Fig. 4 was heated. In Fig. 4, 10 is a roller for conveyance, and the roller with which 11 was water cooled, and 12 is a heat shielding plate. The silicon substrate 8 is heated by the heater 9 installed in the central part of the thermal shield version 12 in the process in which it moves to an arrow direction, and 9'. Although temperature rise of the part for the heater direct lower part of a silicon substrate is carried out, since heat is intercepted with a water cooled roller and the portion which got used from there is maintained at low temperature, firm junction does not take place. Heater power was adjusted so that the silicon substrate temperature directly under a heater might be 550 degrees C in this example.

[0029]Generating of the void was not seen although the silicon substrate joined as mentioned above was observed with the infrared camera like the working example 3. The shearing stress at the time of destruction was 91 kg/cm².

[0030]

[Effect of the Invention] As described above, the effect that the joining method of the present invention enables firm junction of a silicon substrate at low temperature compared with the former is acquired. According to the method of the present invention, since it is joinable at low temperature, the silicon substrates after element formation can be joined and a means effective in formation of spacial configuration objects, such as a micromachine, can be provided.

[0031]Since film peeling by the thermal expansion difference of a laminated structure, generating of a void, etc. can be lessened, quality and reliability can also improve.

(11)Publication number

08-195334

(43)Date of publication of application

30.07.1996

(51)Int.Cl.

H01L 21/02

H01L 21/308

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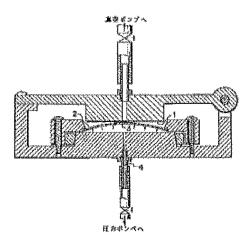
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[Claim 4]A joining method of the silicon substrate according to claim 1 after irradiating inactive ions accelerated to a substrate surface in a high vacuum and forming a pure silicon surface, wherein the aforementioned hydrophobing processing introduces hydrogen gas and carries out the termination of the substrate surface with a hydrogen atom.

[Claim 5]A joining method of the silicon substrate according to claim 1 contacting sequentially a process which the aforementioned substrates are contacted under application of pressure, and is joined from a central part of the aforementioned substrate.

[Claim 6]A joining method of the silicon substrate according to claim 1 performing the aforementioned heat-treatment at 250 degrees C - 700 degrees C.

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A substrate is joined via an oxidizing zone.

[0003]

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[0008]Next, the substrate of two sheets by which hydrophobing processing was carried out is contacted while applying a pressure. Fig.1 is a schematic view of the equipment used for contact of a substrate by the present invention. This equipment holds the first substrate 1 evenly, introduces the fluid (either a liquid or a gas is OK) pressurized from the fluid introducing port 4, and it makes the substrate of two sheets approach, applying this hydrostatic pressure to the back surface of the second substrate 2. Since hydrostatic pressure is applied to the thin membrane 3 and this membrane is bent by the equipment of Fig.1 in sphere form, the 2nd substrate 2 that touches it is also bent by it in sphere form. When the 2nd substrate 2 bends, the substrate of two sheets contacts from each central part. Since a controlled atmosphere is driven out by the periphery in a contact process in this contact method, into the atmosphere, gas does not have a residual thing [carrying out] between substrates. Of course, the substrate of two sheets may be contacted in a vacuum. If, as for hydrostatic pressure, there is more than 3 kg/cm², it is effective enough.

[0009]Next, it heats in order to strengthen junction of the silicon substrate of two sheets which contacted. In the present invention, if the heating temperature is 250 degrees C or more, practically sufficient bonding strength will be obtained, but 300 degrees C - 700 degrees C, if it is 500-700 degrees C still more preferably, the firm junction which is equal to the conventional method will be attained. The temperature rise of the joined substrate is carried out by heating means, such as heater heating and lamp heating. Restriction may not be in junction and any may be sufficient as the atmosphere under heating among the atmosphere, inactive gas, and a vacuum.

[0010] The effect to heat is guessed as follows. When the hydrogen atom which is sticking to a silicon substrate surface by heating disconnects itself, The silicon atoms in the substrate outermost surface couple directly, or a hydrogen atom and the oxygen atom in the atmosphere place and change, and when a silicon atom joins together via an oxygen atom, junction of the silicon substrate of two sheets is performed.

[0011]In the increasing process of the bonding strength under heating, when a strong combination progresses from a substrate periphery, a hole (void) may often occur in the central part of the substrate. It is thought that the portion once joined firmly remains as a void as a result of being unable to ease a strain but a strain concentrating on an unsealed part. Such a void occurs, also when the portion firmly joined by the heterogeneity of substrate temperature is made at random.

[0012] Therefore, in order to join a silicon substrate, without generating a void, joining from one point is effective. It is effective to pinch the silicon substrate 5 which contact completed as the method, by the board 6 and 6' which are two sheets whose thermal conductivity, such as quartz and ceramics, is comparatively small, as shown in Fig.3, and to heat the center of a substrate of these boards in back. The silicon substrate of two sheets is gradually heated from a center by such a means, a firm junction spreads on a periphery from a center, and generating of a void is suppressed.

[0013]As shown in Fig.4 as other methods, the short side direction of the long and slender heating means 9 and 9' is made to convey the contacted silicon substrate 8 with the conveying roller 10, and there is the method of forming a firm junction gradually from the end of a substrate. The shield 12 and the cooling roller 11 serve to control a temperature rise part by making heat conduction to an unsealed part be deteriorated. The temperature gradient of the substrate of two sheets is made as small as possible by 2 sets of upper and lower sides installing a heating means also in which method of Fig.3 and Fig.4. The same effect is expectable even if it moves a heating means from the end of a

substrate instead of conveying a substrate in the method shown in Fig.4.

[0014]

[Function] According to the present invention, in order to carry out hydrophobing processing of the substrate surface beforehand, firm joining becomes possible also for low temperature.

[0015] Hydrophobing processing can be effectively performed by performing the aforementioned hydrophobing processing by fluoric acid and buffer fluoric acid.

[0016] After the aforementioned hydrophobing processing irradiates the inactive ions accelerated to the substrate surface in the high vacuum and forms a pure silicon surface, it can carry out hydrophobing of a substrate surface completely by introducing hydrogen gas and carrying out the termination of the substrate surface with a hydrogen atom.

[0017] By making it contact sequentially from the central part of the aforementioned substrate, it is lost that a base substance etc. are included of the process which the aforementioned substrates are contacted under application of pressure, and is joined in a bonding interface, and it can prevent generating of a void.

[0018]Sufficient bonding strength can be obtained by performing the aforementioned heat-treatment at 250 degrees C - 700 degrees C.

[0019]

[Working example]

[Working example 1] The silicon substrate of two sheets was contacted in the atmosphere, carrying out surface hydrophobing processing of (100) the silicon substrate of two sheets by fluoric acid 2%, and applying 5 atmosphere gas pressure with the contactor of Fig.1 immediately.

[0020] Next, in the atmosphere, it heated for 2 hours and joined. Heating temperature (heat treatment temperature) was made into each temperature shown in Fig.2.

[0021]In this way, the flexural strength of the produced junction sample was measured. Fig.2 illustrates the interface shearing stress at the time of the destruction computed from flexural strength. The disruptive strength of the silicon substrate measured in the similar way was 55 kg/cm², and when heating at 250 degrees C, it turned out that it is joined sufficiently firmly.

[0022] [Working example 2] (100) After carrying out ultraviolet treatment of the silicon substrate for 10 minutes at 200 degrees C and removing the organic impurities of a substrate surface, hydrophobing processing was carried out by the buffer fluoric acid adjusted to PH4 with ammonium fluoride.

[0023] Heat-treatment was performed after contacting the substrate of two sheets like the working example 1 after that.

[0024]When strength was measured like the working example 1, joining by equivalent strength was confirmed.

[0025] [Comparative example] (100) Hydrophilization treatment of the silicon substrate was carried out, when the bonding strength of the sample which carried out contact and heat-treatment like the working example 1 was measured, as shown in Fig.2, above 600 degrees C, it joined firmly, but strength sufficient at low temperature was not obtained. In the case of 400 degrees C or less, it exfoliated in the plane of composition at the time of dicing saw cutting.

[0026][Working example 3] After (100) the silicon substrate of two sheets was immersed for 5 minutes into sulfuric acid and a hydrogen peroxide mixed aqueous solution of 85 degrees C, pure

water washed for 10 minutes. It pulled and continued, and performed and carried out pure water washing of the hydrophobing processing for 2 minutes for 20 seconds in the 2% fluoric acid aqueous solution. It was made to contact, applying 5 atmospheres with the contactor which showed the silicon substrate of two sheets which pretreated more than to Fig. 1. Next, as an outline was shown in Fig. 3, the two alumina ceramic boards 6 and 6' pinched the silicon substrate 5 of two sheets which contacted, and the central part of this alumina ceramic board was heated by the heater 7 and 7' so that heating temperature might be 500 degrees C.

[0027]Generating of the void was not seen, when infrared rays were irradiated to the silicon substrate of two sheets joined as mentioned above and the transmitted light was observed with the infrared camera. The shearing stress at the time of destruction is 85 kg/cm², and having sufficient strength was checked.

[0028][Working example 4] After irradiating ultraviolet rays for 20 minutes, heating (100) the silicon substrate of two sheets at 200 degrees C in oxygen environment, pure water washing of the hydrophobing processing was performed and carried out for 2 minutes for 20 seconds in the 2% fluoric acid aqueous solution. It was made to contact, applying 5 atmospheres with the contactor which showed the silicon substrate of two sheets which performed the above hydrophobing processing to Fig.1. Next, the silicon substrate 8 which contacted with the heating apparatus which has a sample conveying machine which shows an outline to Fig.4 was heated. In Fig.4, 10 is a roller for conveyance, and the roller with which 11 was water-cooled, and 12 is a heat shielding plate. The silicon substrate 8 is heated by the heater 9 installed in the central part of the thermal shield version 12 in the process in which it moves to an arrow direction, and 9'. Although temperature rise of the part for the heater direct lower part of a silicon substrate is carried out, since heat is intercepted with a water-cooled roller and the portion which got used from there is maintained at low temperature, firm junction does not take place. Heater power was adjusted so that the silicon substrate temperature directly under a heater might be 550 degrees C in this example.

[0029]Generating of the void was not seen although the silicon substrate joined as mentioned above was observed with the infrared camera like the working example 3. The shearing stress at the time of destruction was 91 kg/cm².

[0030]

[Effect of the Invention] As described above, the effect that the joining method of the present invention enables firm junction of a silicon substrate at low temperature compared with the former is acquired. According to the method of the present invention, since it is joinable at low temperature, the silicon substrates after element formation can be joined and a means effective in formation of spacial configuration objects, such as a micromachine, can be provided.

[0031]Since film peeling by the thermal expansion difference of a laminated structure, generating of a void, etc. can be lessened, quality and reliability can also improve.

公告本

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	發謝	· 明 專 利 説 明 書
一、 發明 一、	中文	以三氟化氮退火處理製程改 善轉種氯化層可靠度之方法
新型	英文	
	姓名	(1) 趙 天 生 (2) 雷 添 福 (3) 張 子 云 (4) 周 宗 賢 (5) 陳 希 寛
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本紙張尺度適用中國國家標準 (CNS) A4規格 (210×297公釐)
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SAMSUNG EXHIBIT 1002 Samsung v. Invenssas Bonding

訂

五、發明説明(1)

[發明之範圍]

本發明係關於一種改善剛極氧化層可靠度的方法,特別是利用三氟化氮(NF3)對閘極退火處理的製程,以降低閘極漏腦流的方法。

[發明之背景]

隨著半導體元件的縮小(scale down),關極氧化層(gate oxide)的可靠度也變得更為重要,如何獲得高可靠度的關極氧化層就變成當今最重要的問題。關極(gate)漏電流(leakage current)是一個判斷關極氧化層優劣的主要因素,尤其是在FN穿透(F-N tunneling)前低電場下的漏電流。因此,如何得到一個低漏電流的關極氧化層便成為當今主要的問題。

目前有幾項技術可增進剛極氧化層的品質,例如,在形成氧化層的同時通入少量的氣(F),使其與氧氣同時在砂晶圖上反應:又如,使用氮離子佈植(ion implantation)在多晶砂閘極(poly-Si-gate),對於閘極氧化層的品質也有所改善。

而就另一方面來說,在深次微米的製程中,如何減低閘極的電阻也是一個重要的關鍵。傳統的鈦砂化物(Ti-silicide)雖可以有效的降低閘極電阻,但其在閘極線寬變窄後,其電阻值也會隨之而變大。鈷砂化物(Co-silicide)是一個取代的方法,因為鈷砂化物的電阻值不會因線寬的變窄而變大;但鈷砂化物又存在著另一個問題,其漏電流太大。

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訂

五、發明説明(2)

〔發明之目的與概述〕

本發明之目的在於,提供一種改善關極氧化層可靠度的方法,特別是利用三氟化氮 (NFa)對關極氧化層做退火處理的製程,其可顯著改善關極氧化層的漏電流及崩潰電場。本發明之另一目的在於,提供一種改善關極氧化層可靠度的方法,特別是利用三氟化氮 (NFa)對關極氧化層做退火處理的製程,其可有效減少因形成關極鈷砂化物而來的漏電流。

本發明之以三氟化氮退火處理製程改善關極氧化層可靠度之方法,係一項結合氮、氟優點的技術來改善關極氧化層的可靠度,其係以三氟化氮(NFa)對多晶砂(poly-Si)關極作退火(annealing)處理,使氟化砂(Si-F)與氮化砂(Si-N)的鍵結在二氧化砂與矽晶圓的界面(SiOz/Siinterface)中產生,此種較強的鍵結會取代較弱的鍵結,例如氫化砂(Si-H)與氫氧化砂(Si-OH),而使得氧化層的可靠度增強。

[附圖之簡要說明]

圖 1係為本發明製程之一實施例之試片製作流程圖。 圖 2係為不同退火時間的鋁電極試片之崩潰電場分佈圖。 圖 3係為不同退火時間的鋁電極試片之漏電流分佈圖。 圖 4係為以三氟化氮退火處理試片之二次離子分佈圖。 圖 5條為本發明製程之另一實施例之試片製作流程圖。 圖 6條為不同退火時間的鈷砂化物電極試片之崩潰電場

分佈圖。

五、發明説明(3)

圖 7係為不同退火時間的鈷砂化物電極試片之漏電流分佈圖。

[發明之詳細說明]

以下參照附圖說明本發明之實施例。

如圖1所示本發明製程之一實施例之試片製作流程圖,在P型(100)矽晶圖上製作一金氧半電容(MOS capacitor),其步驟為:首先,以傳統爐管在晶圖上形成一層 5000埃的阻隔氣化層 (field oxide),並蝕刻出一主動區 (active region)。其次,在用氦氣稀釋的氦氣下(Nz:02=10:1),形成一約 33埃厚的超薄氧化層 (ultra-thin oxide)(該超薄氧化層的厚度並不限定於 33埃,其範圖可為 15至 70埃),並用低壓化學氣相沉積法 (LPCVD)堆疊一層 3000埃的多晶砂 (poly-Si)。然後,在攝氏 620度低壓的環境下,用三氟化氮 (NFa)對多晶砂做不同時間 (5至 20分鐘)的退火處理。最後,以鉛做成閘極電極。

在上述以三氟化氮對多晶的做退火處理的過程中,會有氟化砂(Si-F)與氮化矽(Si-N)的鍵結在二氧化矽與矽晶圓的界面(SiOz/Si interface)中產生,此種較強的鍵結會取代較弱的鍵結,例如氫化矽(Si-H)與氫氧化矽(Si-OH),而使得氧化層的可靠度增強。如圖4所示,經由對三氟化氮退火處理的試片做二次離子分析(SINS),可確知在二氧化矽與矽晶圓的界面有氟與氦的鍵結形成。

圖 2 及 圖 3 分 別 為 不 同 退 火 時 間 之 鋁 電 極 試 月 的 崩 潰 電 場 (breakdown field)及 漏 電 流 (leakage current)之 統 計 分

訂

五、發明説明(4)

佈圖 , 其 中 使 用 三 氟 化 氮 做 退 火 處 理 的 試 片 其 崩 潰 電 場 皆 比 對 照 組 試 月 高 , 且 其 漏 電 流 亦 皆 比 對 照 組 試 月 低 。

如圖 5 所示本發明製程之另一實施例之試片製作流程圖,其中,除了最後步驟係以鈷化矽做成剛極電極外,其餘步驟時與前一實施例相同。圖 6 及圖 7分別為不同退火時間之鈷化矽電極試片的崩潰電場及漏電流之統計分佈圖,其中使用三氟化氮做退火處理的試片其崩潰電場皆比對照組試片低。又,在此實施例中,以三氟化氮對鈷化矽電極試片做退火處理的時間之嚴佳化條件係為 10分鐘。

綜上所述,以三氟化氮對多晶矽做退火處理,確可顯著 改善氧化層的崩潰電場及漏電流,增進氧化層的可靠度。 本發明雖已參照附屬及實施例詳細說明,然其並不囿限 於上述二實施例。任何依本發明之精神及申請專利範圍所 做的變化及修改,仍應屬於本發明之範圍。

)

以三氟化氮退火處理製程改善閘極⁾ 氧化層可靠度之方法

本發明之以三氟化氮退火處理製程改善關極氧化層可靠度之方法,係一項結合氮、氟優點的技術來改善關極氧化層的可靠度,其係以三氟化氮(NF3)對多晶矽(poly-Si)關極作退火(annealing)處理,使氟化矽(Si-F)與氮化矽(Si-N)的鍵結在二氧化矽與矽晶圓的界面(SiOz/Siinterface)中產生,此種較強的鍵結會取代較弱的鍵結,例如氫化矽(Si-H)與氫氧化砂(Si-OH),而使得氧化層的可靠度增強。

英文發明摘要(發明之名稱:

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六、申請專利範圍

1.一種以三氟化氮退火處理製程改善閘極氧化層可靠度之方法,其具有下列主要步驟:

在晶圓上形成一阻隔氧化層(field oxide),並蝕刻一主動區(active region);

形成一超薄氧化層(ultra-thin oxide), 並堆疊一層多 品砂(poly-Si);

以三氟化氮 (NF3) 對多晶矽作退火處裡;及 形成關極電極。

- 2.如申請專利範圍第1項之方法,其中該超薄氧化層之厚度係為15至70埃。
- 3.如申請專利範圍第1項之方法,其中該多晶砂係以低 壓化學氣相沉積(LPCVD)堆疊。
- 4.如申請專利範圍第1項之方法,其中該退火處理係於攝氏620度低壓的環境下進行。
- 5.如申請專利範團第1項之方法・其中該退火處理之時間係為5至20分鐘。
- 6. 如申讀專利範團第1項之方法,其中該關極電極係為鋁電極。
- 7. 如申請專利範圖第1項之方法,其中該層極電極係為

 鈷砂化物電極。
- 8. 如申請專利範圍第7項之方法,其中該退火處理之時間係以10分鐘為最佳。

在P型(100) 矽晶片上形成阻隔氧化層(field oxide)
於稀釋O2下形成33埃之超薄氧化層(ultra-thin oxide)

堆疊一層3000埃之多晶矽(poly-Si),隨後摻雜POCl,
以NF,對多晶矽作不同時間的退火處理

形成Al電極

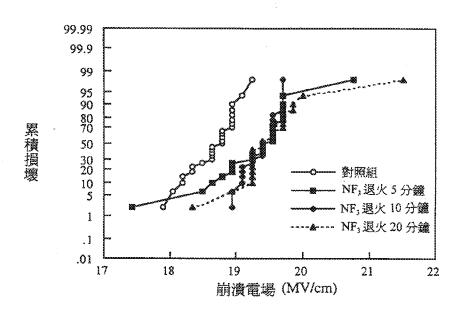
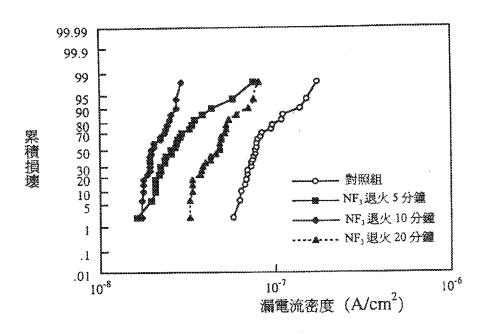
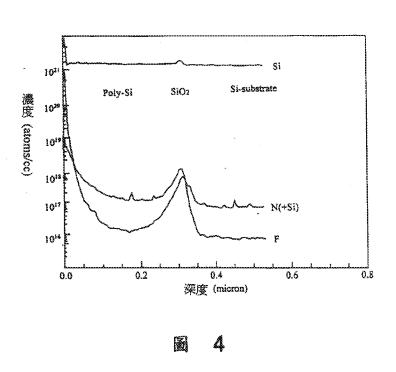


图 2





在 P型 (100) 矽晶片上形成阻隔氧化層 (field oxide)

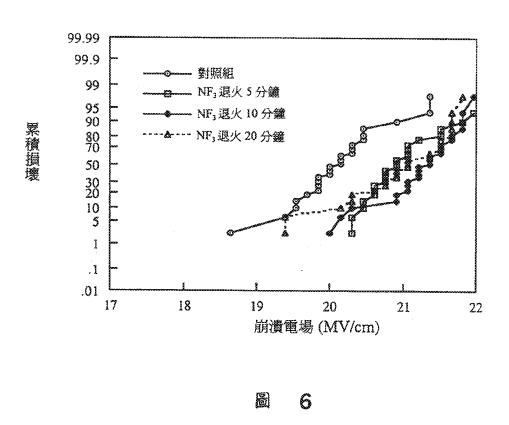
於稀釋 O₂下形成 33 埃之超薄氧化層 (ultra-thin oxide)

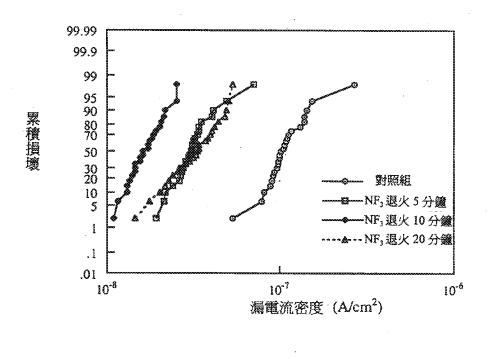
堆疊一層 3000 埃之多晶矽 (poly-Si),隨後摻雜 POCl₃

以 NF₃對多晶矽作
不同時間的退火處理

以二階段退火 (two-step annealing) 形成 CoSi₂電極

圖





1 TW389965B 2000-05-11 Method for improving reliability of a gate oxide layer by NF3 annealing (en) 2 (Electronic English Translation) 3 **English Abstract:** 4 5 6 This invention provides a method for improving reliability of a gate oxide layer by NF3 annealing, which 7 combines the advantages of nitrogen and fluorine to enhance the reliability of the gate oxide layer. A 8 polysilicon gate is annealed by using NF3 to produce Si-F and Si-N bonding at the Si/SiO2 interface. Such 9 stronger bonding replaces the weaker bonding such as Si-H and Si-OH to enhance the reliability of the 10 oxide layer. 11 12 **English Description:** 13 14 15 Technical Field 16 17 This invention is about a kind of improved gate method for reliable degree of the oxide layer, in 18 particular through the use of nitrogen trifluoride (NF 3) the manufacturing process of treatment of the 19 gate annealing, in order to reduce the gate leakage current method. 20 21 Background Art 22 23 Reduced along with the semiconductor element (scale down), the reliability of the gate oxide layer (gate 24 oxide) has also become more important, how to obtain high-reliability of the gate oxide layer to become 25 the most important issue. Gate (gate) leakage current (leakage current) is a judging gate oxide layer of 26 the main factors, especially in the low electric field before penetrating FN (F-N tunneling) drain current. 27 Therefore, how to get a low leakage current of the gate oxide layer will become a major problem today. 28 29 There are currently several technology can improve the quality of the gate oxide layer, for example, at 30 the same time forming the oxide layer of the inject a small amount of fluorine (F), the oxygen reaction 31 on a silicon wafer at the same time; and if, the polycrystalline (ion implantation) using nitrogen ion 32 implanter is [...] (poly-Si-gate), the quality of the gate oxide layer has also improved. 33 34 On the other hand, in the deep sub-micron in the process of, how to reduce gate resistance is also an 35 important key. Although traditional [...] compound (Ti-silicide) gate resistance can be reduced 36 effectively, but the narrowing of the line width of the gate, its resistance will be thereupon and becomes 37 larger. (Co-silicide) cobalt silicon compound is a substituted method, because the resistance of the 38 cobalt silicon compound will not become great due to the narrowing of the line width; but the cobalt 39 silicon compound also exists another problem works, its leakage current is too large.

40	
41	Summary of the Invention
42	
43	The purpose of this invention is to, provide a degree of improved gate method of the oxide layer, in
44	particular through the use of nitrogen trifluoride (NF 3) gate oxidation layer as the process of annealing,
45	which can remarkably improve the leakage current of the gate oxide layer and collapse the electric field.
46	
47	Another purpose of this invention is to, provide a degree of improved gate method of the oxide layer, in
48	particular through the use of nitrogen trifluoride (NF 3) gate oxidation layer as the process of annealing,
49 50	the formed can be effectively reduced to cobalt silicon compound gate leakage current.
51	In the invention nitrogen trifluoride gate oxide layer to improve the process for preparing of annealing
52	treatment method for reliability, is a combined nitrogen, fluoro the advantages of the technology to
53	improve the reliability of the gate oxide layer, the nitrogen trifluoride (NF 3) of the polycrystalline silicon
54	(poly-Si) gate as the annealing treatment (annealing), the silicon nitride silicon fluoride with (Si-F) (Si-N)
55	the bonding of the silicon wafer, the silicon dioxide interface (SiO 2/Si interface) generated in, such
56	strong the key ties the meeting substituted weaker bonding, for example with silicon hydroxide (Si-H)
57	(Si-OH) [] , enhanced reliability of the oxide layer.
58	
59	Description of Drawings
60	
61	Figure 1 is the embodiment of the process of this invention one of the manufacturing flow chart of the
62	test piece.
63	
64	Figure 2 is a different annealing time collapse of the aluminum electrode of the electric field distribution
65	map of the test piece.
66	
67	Figure 3 is a different annealing time aluminum electrode of the leakage current distribution map of the
68	test piece.
69 	
70	Figure 4 is to nitrogen trifluoride annealing treatment test piece secondary ion distribution map of.
71	
72 73	Figure 5 is the manufacturing process for this invention another embodiment of the manufacturing flow
73 74	chart of the test piece.
74 75	Figure 6 is a different annealing time of the test piece of the electrode of the cobalt silicon compound
75 76	collapse of the distribution of the electric field.
70 77	collapse of the distribution of the electric field.
77 78	Fig. 7 is a different annealing time of the cobalt silicon compound electrode leakage current distribution
70 79	map of the test piece.
79 80	map of the test piece.
81	(Detailed description of invention)

By the following with reference to the embodiment of the present invention.

As shown in Figure 1 one of the invention the embodiment of the test process of the manufacturing flow chart, in the P-type (100) silicon wafer (MOS capacitor) making a MOS capacitor, its steps are: first, to traditional furnace tube is formed on the wafer a layer of 5000 angstroms (field oxide) of the barrier oxide layer, and etching the active region (active region). Secondly, in a nitrogen-diluted oxygen (N 2: O 2 = 10:1), forming a approximately 33 angstroms thick ultra-thin oxide layer (ultra-thin oxide) (the ultra-thin the thickness of the oxide layer does not defines in 33 heydon, its range can be is 15 to 70 angstroms), and using low pressure chemical vapor deposition (LPCVD) stacking a layer of 3000 angstroms of polycrystal silicon (poly-Si). Furthermore, at 620 degrees under low-pressure environment, nitrogen trifluoride for (NF 3) of the polycrystalline silicon to do different time (5 to 20 minutes) the annealing treatment. Finally, the gate electrode is made of aluminum.

In the above-mentioned by using nitrogen trifluoride of the polycrystalline silicon as in the process of annealing, with a silicon nitride silicon fluoride (Si-N) (Si-F) the bonding of the silicon wafer, the silicon dioxide interface (SiO 2/Si interface) generated in, such strong the key ties the meeting substituted weaker bonding, for example with silicon hydroxide (Si-H) (Si-OH) [...], enhanced reliability of the oxide layer. As shown in Figure 4 is shown, through the nitrogen trifluoride annealing treatment of the test piece to the secondary ion analysis (SIMS), can be sure in the silicon dioxide with the interface of the silicon wafer with the nitrogen to form a bonded fluorine.

Figure 2 and Figure 3 are respectively for different annealing time of the collapse of the aluminum test piece of the electrode and the drain current (leakage current) electric field (breakdown field) the statistical profile, wherein to use nitrogen trifluoride its collapse test piece annealing treatment of the electric field than the control test piece are high, and the leakage current test than the control group are also low.

As shown in Figure 5 the invention the embodiment of the process for preparing the test piece preparation flowdiagram, wherein the final steps in addition to the gate electrode silicon cobalt is made, the remaining steps are the same as the previous embodiment. Figure 6 and Figure 7 are respectively for different annealing time of the collapse of the cobalt [...] the test piece of the electrode of the electric field and leakage current statistical profile, wherein to use nitrogen trifluoride its collapse test piece annealing treatment of the electric field than the control test piece are high, and the leakage current test than the control group are also low. Also, in this embodiment, the nitrogen trifluoride cobalt in silicon the test piece of the electrode to the time of annealing to the optimization conditions for 10 minutes.

To sum up, by using nitrogen trifluoride of the polycrystalline silicon as annealing processing, the oxide layer can remarkably improve the collapse of the electric field and leakage current, to enhance the reliability of the oxide layer.

124	The invention, although reference has been made to the attached drawing and detailed description of
125	the embodiment, however, it does not the park is restricted in the above-mentioned second
126	embodiment. Any in accordance with this invention the spirit and the scope of the Patent application
127	the change and modification, should still belongs to the scope of this invention.
128	
129	ENGLISH CLAIMS:
130	
131	
132	A nitrogen trifluoride gate oxide layer to improve the process for preparing of annealing treatment
133	method for reliability, it has the following main steps: on the wafer (field oxide) forming a barrier oxide
L34	layer, and etching a active region (active region) ; (ultra-thin oxide) to form a ultra-thin oxide layer, and
L35	stacking a layer of polycrystal silicon (poly-Si); to nitrogen trifluoride (NF 3) of the polycrystalline silicon
136	for annealing process; and forming the gate electrode.
137	
138	Method according to Claim 1, wherein the ultra-thin oxide layer of a thickness of 15 to 70 angstroms.
139	
140	Method according to Claim 1, wherein the polycrystalline silicon is stacked low-pressure chemical vapor
141	deposition (LPCVD).
142	
143	Method according to Claim 1, wherein on the annealing treatment at 620 degrees the low-pressure
144	environment.
145	
146	Method according to Claim 1, wherein the annealing treatment time is 5 to 20 minutes.
147	
148	Method according to Claim 1, wherein the gate electrode is an aluminum electrode.
149	
150	Method according to Claim 1, wherein the gate electrode is a cobalt silicon compound electrode.
151	

Method according to Claim 7, wherein the annealing treatment uses the time of 10 minutes is the best.

Electronic Ack	knowledgement Receipt
EFS ID:	24585269
Application Number:	14957501
International Application Number:	
Confirmation Number:	1033
Title of Invention:	METHOD FOR LOW TEMPERATURE BONDING AND BONDED STRUCTURE
First Named Inventor/Applicant Name:	Qin-Yi Tong
Customer Number:	135980
Filer:	Paul Steven Stellman/Jonathan Pan
Filer Authorized By:	Paul Steven Stellman
Attorney Docket Number:	TSSRA.012C1
Receipt Date:	11-JAN-2016
Filing Date:	02-DEC-2015
Time Stamp:	16:05:39
Application Type:	Utility under 35 USC 111(a)

Payment information:

Submitted with I	Payment		no			
File Listing:						
Document Number	Document Description		File Name	File Size(Bytes)/ Message Digest	Multi Part /.zip	Pages (if appl.)
1		2010	6-01-11_IDS_TSSRA-012C1.	166178	yes	7
·			pdf	32cffa4a9d5744dedb5e4fb402e69026db8 5c443	· · · · · · · · · · · · · · · · · · ·	•

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	Transmittal	Letter	1	2			
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Warnings:							
Information:							
2	Foreign Reference	CA2399282A1.pdf	1664146	no	49		
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3	Foreign Reference	JP-H08-195334.pdf	1116815	no	17		
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51	Non Patent Literature	bonded-SOI-bipolar-transistor. pdf	6ea83585af9b78b1d4318a76d50270c59fe 33f04	no	10
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47	Non Patent Literature	Roberds-et-al_Low- temperature-in-situ-plasma-	3260176	no	9
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45	Non Patent Literature	Reiche-et-al_The-effect-of-a-	4357119	no	8

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New Applications Under 35 U.S.C. 111

If a new application is being filed and the application includes the necessary components for a filing date (see 37 CFR 1.53(b)-(d) and MPEP 506), a Filing Receipt (37 CFR 1.54) will be issued in due course and the date shown on this Acknowledgement Receipt will establish the filing date of the application.

National Stage of an International Application under 35 U.S.C. 371

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New International Application Filed with the USPTO as a Receiving Office

If a new international application is being filed and the international application includes the necessary components for an international filing date (see PCT Article 11 and MPEP 1810), a Notification of the International Application Number and of the International Filing Date (Form PCT/RO/105) will be issued in due course, subject to prescriptions concerning national security, and the date shown on this Acknowledgement Receipt will establish the international filing date of the application.

Docket No.: TSSRA.012C1 Customer No. 135,980

INFORMATION DISCLOSURE STATEMENT

Inventor : Tong et al.

App. No. : 14/957,501

Filed : December 2, 2015

For : METHOD FOR LOW

TEMPERATURE BONDING AND

BONDED STRUCTURE

Examiner : Unknown

Art Unit : 2895

Conf. No. : 1033

Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450

References and Listing

Pursuant to 37 CFR 1.56, an Information Disclosure Statement listing references is provided herewith. Copies of any listed foreign and non-patent literature references are being submitted.

No Disclaimers

To the extent that anything in the Information Disclosure Statement or the listed references could be construed as a disclaimer of any subject matter supported by the present application, Applicant hereby rescinds and retracts such disclaimer.

Timing of Disclosure

This Information Disclosure Statement is being filed within three months of the filing date, and no fee is believed to be required.

The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment, to Account No. 11-1410.

Application No.: 14/957,501

Filing Date: December 2, 2015

Respectfully submitted,

KNOBBE, MARTENS, OLSON & BEAR, LLP

Dated: January 11, 2016 By: /Paul Stellman/

Paul Stellman Registration No. 68,287 Attorney of Record Customer No. 135,980 (415) 954-4114

Docket No.: TSSRA.012C1 January 11, 2016
Page 1 of 1

Please Direct All Correspondence to Customer Number 135,980

EFS WEB CONTINUING IDS COVER LETTER

Inventor : Tong et al.

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Examiner : Unknown

Art Unit : 2895

Conf No. : 1033

Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450

Dear Sir:

Submitted herewith are references numbered 55 to 66 listed on the PTO/SB/08 or equivalent filed under EFS ID 24585269.

Please charge any additional fees, including any fees for additional extension of time, or credit overpayment to Deposit Account No. 11-1410.

Respectfully submitted,

KNOBBE, MARTENS, OLSON & BEAR, LLP

Dated: January 11, 2016 By: /Paul Stellman/

Paul Stellman Registration No. 68,287 Attorney of Record Customer No. 135,980 (415) 954-4114

(413) 934-41

Electronic Acknowledgement Receipt				
EFS ID:	24587990			
Application Number:	14957501			
International Application Number:				
Confirmation Number:	1033			
Title of Invention:	METHOD FOR LOW TEMPERATURE BONDING AND BONDED STRUCTURE			
First Named Inventor/Applicant Name:	Qin-Yi Tong			
Customer Number:	135980			
Filer:	Paul Steven Stellman/Jonathan Pan			
Filer Authorized By:	Paul Steven Stellman			
Attorney Docket Number:	TSSRA.012C1			
Receipt Date:	11-JAN-2016			
Filing Date:	02-DEC-2015			
Time Stamp:	16:13:27			
Application Type:	Utility under 35 USC 111(a)			

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1	Transmittal Letter	2016-01-11_Cont- IDS_TSSRA-012C1.pdf	63258	no	1
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Page 178 ion 331

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Warnings:			0e93eb65b1aa2bf50364b7e165e5ac48e2a c1f08		
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2	Non Patent Literature	Shajii-et-al_A-backside- contact-technology.pdf	3174356	no	9

11	Non Patent Literature	Tong-et-al_Wafer-bonding-	950230	no	17
11	Non ratent Literature	and-layer-splitting.pdf	c1cc801e40d43b3fac829d37686ffc4676b0 e053		''
Warnings:					
Information:					
12	Non Patent Literature	Weldon-et-al_Physics-and- chemistry-of-silicon-wafer-	748476	no	13
12	World attended to	bonding.pdf	0c02a3e27f61b0091e2e3e5f8dcb6da88eb 107df		
Warnings:					
Information:					
13	Non Patent Literature	Yallup_Analog-CMOS-circuits- on-thick-films.pdf	4201931	no	13
			290b9043e310805baf3f24e373f2e2cb06fd 3804	110	
Warnings:					
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		Total Files Size (in bytes)	286	511743	

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If a new application is being filed and the application includes the necessary components for a filing date (see 37 CFR 1.53(b)-(d) and MPEP 506), a Filing Receipt (37 CFR 1.54) will be issued in due course and the date shown on this Acknowledgement Receipt will establish the filing date of the application.

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If a timely submission to enter the national stage of an international application is compliant with the conditions of 35 U.S.C. 371 and other applicable requirements a Form PCT/DO/EO/903 indicating acceptance of the application as a national stage submission under 35 U.S.C. 371 will be issued in addition to the Filing Receipt, in due course.

New International Application Filed with the USPTO as a Receiving Office

If a new international application is being filed and the international application includes the necessary components for an international filing date (see PCT Article 11 and MPEP 1810), a Notification of the International Application Number and of the International Filing Date (Form PCT/RO/105) will be issued in due course, subject to prescriptions concerning national security, and the date shown on this Acknowledgement Receipt will establish the international filing date of the application.



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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
14/957,501	12/02/2015	Qin-Yi Tong	TSSRA.012C1	1033	
135980 Knobbe, Marte i	7590 02/03/201 ns. Olson	6	EXAM	IINER	
& Bear, LLP 2040 main Stree			GARCIA, J	OANNIE A	
Fourteenth Floo			ART UNIT	PAPER NUMBER	
Irvine, CA 926	14		2895		
			NOTIFICATION DATE	DELIVERY MODE	
			02/03/2016	ELECTRONIC	

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Notice of the Office communication was sent electronically on above-indicated "Notification Date" to the following e-mail address(es):

jayna.cartee@knobbe.com efiling@knobbe.com

		Application No.Applicant(s)14/957,501TONG ET AL			
	Office Action Summary	Examiner JOANNIE A. GARCIA	Art Unit 2895	AIA (First Inventor to File) Status No	
T	The MAILING DATE of this communication app Reply	ears on the cover sheet with the	correspondence	e address	
A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTHS FROM THE MAILING DATE OF THIS COMMUNICATION. - Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication. - If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication. - Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).					
Status					
	esponsive to communication(s) filed on <u>Dece.</u> declaration(s)/affidavit(s) under 37 CFR 1.1 :				
2a) 🗌 Th	is action is FINAL . 2b)⊠ This	action is non-final.			
3) □ A n	election was made by the applicant in respo	onse to a restriction requiremen	t set forth durin	g the interview on	
4) Sir	; the restriction requirement and election nee this application is in condition for allowan used in accordance with the practice under E	ce except for formal matters, p	rosecution as to	o the merits is	
Disposition	of Claims*				
5) Claim(s) 2-34 is/are pending in the application. 5a) Of the above claim(s) is/are withdrawn from consideration. 6) Claim(s) is/are allowed. 7) Claim(s) 2-34 is/are rejected. 8) Claim(s) is/are objected to. 9) Claim(s) are subject to restriction and/or election requirement. * If any claims have been determined allowable, you may be eligible to benefit from the Patent Prosecution Highway program at a participating intellectual property office for the corresponding application. For more information, please see http://www.uspto.gov/patents/init_events/pph/index.jsp or send an inquiry to PPHfeedback@uspto.gov.					
Application Papers 10) ☐ The specification is objected to by the Examiner. 11) ☒ The drawing(s) filed on <u>December 2, 2015</u> is/are: a) ☒ accepted or b) ☐ objected to by the Examiner. Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a). Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).					
Priority under 35 U.S.C. § 119 12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f). Certified copies: a) All b) Some** c) None of the: 1. Certified copies of the priority documents have been received. 2. Certified copies of the priority documents have been received in Application No. 3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)). ** See the attached detailed Office action for a list of the certified copies not received.					
Attachment(s)					
1) Notice of	References Cited (PTO-892)	3) Interview Summa			
2) Information Disclosure Statement(s) (PTO/SB/08a and/or PTO/SB/08b) Paper No(s)/Mail Date <u>January 11, 2016</u> . Paper No(s)/Mail Date 4) Other:					

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The present application is being examined under the pre-AIA first to invent provisions.

DETAILED ACTION

This action is responsive to: the Application filed December 2, 2015, and the Information Disclosure Statement (IDS) Document, Form PTO-1449, filed January 11, 2016.

Claims 2-34 are pending. Claims 1, 22, 32, and 33, are independent.

Information Disclosure Statement

Acknowledgment is made of applicant's Information Disclosure Statement (IDS)

Document filed January 11, 2016.

This Information Disclosure Statement (IDS) Document has been considered, and a signed and dated copy is included herein.

Double Patenting

The nonstatutory double patenting rejection is based on a judicially created doctrine grounded in public policy (a policy reflected in the statute) so as to prevent the unjustified or improper time wise extension of the "right to exclude" granted by a patent and to prevent possible harassment by multiple assignees. A nonstatutory obviousness-type double patenting rejection is appropriate where the conflicting claims are not identical, but at least one examined application claim is not patentably distinct

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from the reference claim(s) because the examined application claim is either anticipated by, or would have been obvious over, the reference claim(s). See, e.g., *In re Berg*, 140 F.3d 1428, 46 USPQ2d 1226 (Fed. Cir. 1998); *In re Goodman*, 11 F.3d 1046, 29 USPQ2d 2010 (Fed. Cir. 1993); *In re Longi*, 759 F.2d 887, 225 USPQ 645 (Fed. Cir. 1985); *In re Van Ornum*, 686 F.2d 937, 214 USPQ 761 (CCPA 1982); *In re Vogel*, 422 F.2d 438, 164 USPQ 619 (CCPA 1970); and *In re Thorington*, 418 F.2d 528, 163 USPQ 644 (CCPA 1969).

A timely filed terminal disclaimer in compliance with 37 CFR 1.321(c) or 1.321(d) may be used to overcome an actual or provisional rejection based on a nonstatutory double patenting ground provided the conflicting application or patent either is shown to be commonly owned with this application, or claims an invention made as a result of activities undertaken within the scope of a joint research agreement.

Effective January 1, 1994, a registered attorney or agent of record may sign a terminal disclaimer. A terminal disclaimer signed by the assignee must fully comply with 37 CFR 3.73(b).

Claims 22-31 are rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over claims 1 and 6, of U.S. Patent Application 2015/0303263. Although the conflicting claims are not identical, it is not patentably distinct from each other because the claims of the current application, U.S. Patent Application 14/957,501, are broader than the claims of the U.S. Patent Application 2015/0303263. In this regard, the limitations of instant application claim 22 appear to be anticipated or obvious over '263 claims 1 and 6.

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For example, instant application claim 22 recites, "...a first semiconductor material comprising a planarized insulating material having a first bonding surface; and a second material having a second bonding surface, wherein the first bonding surface is an etched surface terminated with a nitrogen-containing species, and wherein the first and second bonding surfaces are in direct contact with each other and bonded together with a chemical bond without any intervening adhesive...", which includes the similarly recited limitations in '263 claims 1 and 6, namely, "...a processed semiconductor device wafer comprising a first surface and a planarized insulating material having a first bonding surface wherein said planarized insulating material is opposite said first surface; a first material having a second surface and a second bonding surface opposite said second surface, wherein no devices are disposed in said first material; wherein said first bonding surface is an etched surface terminated with a species; and wherein said first and second bonding surfaces are in direct contact with each other and bonded together with a chemical bond without any adhesive...", and "...and wherein said species comprises nitrogen...".

Claims 22-31 are rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over claim 30, of U.S. Patent Application 2015/0303263. Although the conflicting claims are not identical, it is not patentably distinct from each other because the claims of the current application, U.S. Patent Application 14/957,501, are broader than the claims of the U.S. Patent Application

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2015/0303263. In this regard, the limitations of instant application claims 22 and 27 appear to be anticipated or obvious over '263 claim 30.

For example, instant application claim 22 recites, "...a first semiconductor material comprising a planarized insulating material having a first bonding surface; and a second material having a second bonding surface, wherein the first bonding surface is an etched surface terminated with a nitrogen-containing species, and wherein the first and second bonding surfaces are in direct contact with each other and bonded together with a chemical bond without any intervening adhesive...", and "...wherein said planarized insulating material comprises silicon oxide...", which includes the similarly recited limitations in '263 claim 30, namely, "...a processed semiconductor device wafer comprising a first surface having a thickness, a planarized insulating material of deposited silicon oxide having a first bonding surface, wherein said planarized insulating material is opposite said first surface, and a device layer between said first surface and said planarized insulating material; a first material comprising silicon having a second surface and a second bonding surface opposite said second surface, wherein no devices are disposed in said first material; wherein said first bonding surface is an etched surface terminated with nitrogen; wherein said first and second bonding surfaces are in direct contact with each other and bonded together with a chemical bond without any adhesive...".

Claims 22-31 are rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over claims 31 and 36, of U.S. Patent

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Application 2015/0303263. Although the conflicting claims are not identical, it is not patentably distinct from each other because the claims of the current application, U.S. Patent Application 14/957,501, are broader than the claims of the U.S. Patent Application 2015/0303263. In this regard, the limitations of instant application claim 22 appear to be anticipated or obvious over '263 claims 31 and 36.

For example, instant application claim 22 recites, "...a first semiconductor material comprising a planarized insulating material having a first bonding surface; and a second material having a second bonding surface, wherein the first bonding surface is an etched surface terminated with a nitrogen-containing species, and wherein the first and second bonding surfaces are in direct contact with each other and bonded together with a chemical bond without any intervening adhesive...", which includes the similarly recited limitations in '263 claims 31 and 36, namely, "...a first silicon material comprising a first surface and a planarized insulating material having a first bonding surface wherein said planarized insulating material is opposite said first surface; a second material having a second surface and a second bonding surface opposite said second surface, wherein no devices are disposed in said second material; wherein said first bonding surface is an etched surface terminated with a species; and wherein said first and second bonding surfaces are in direct contact with each other and bonded together with a chemical bond without any adhesive..." and "...wherein said species comprises nitrogen...".

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Claims 22-31 are rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over claim 60, of U.S. Patent Application 2015/0303263. Although the conflicting claims are not identical, it is not patentably distinct from each other because the claims of the current application, U.S. Patent Application 14/957,501, are broader than the claims of the U.S. Patent Application 2015/0303263. In this regard, the limitations of instant application claims 22 and 27 appear to be anticipated or obvious over '263 claim 60.

For example, instant application claim 22 recites, "...a first semiconductor material comprising a planarized insulating material having a first bonding surface; and a second material having a second bonding surface, wherein the first bonding surface is an etched surface terminated with a nitrogen- containing species, and wherein the first and second bonding surfaces are in direct contact with each other and bonded together with a chemical bond without any intervening adhesive...", and "...wherein said planarized insulating material comprises silicon oxide...", which includes the similarly recited limitations in '263 claim 60, *namely*, "...a planarized insulating material of deposited silicon oxide having a first bonding surface, wherein said planarized insulating material is opposite said first surface, and a device layer between said first surface and said planarized insulating material; a second material comprising silicon having a second surface and a second bonding surface opposite said second surface, wherein no devices are disposed in said second material; wherein said first bonding surface is an etched surface terminated with nitrogen; wherein said first and second bonding

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surfaces are in direct contact with each other and bonded together with a chemical bond without any adhesive...".

Claims 22-31 are rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over claims 77 and 82, of U.S. Patent 8,153,505. Although the conflicting claims are not identical, it is not patentably distinct from each other because the claims of the current application, U.S. Patent Application 14/957,501, are broader than the claims of the U.S. Patent 8,153,505. In this regard, the limitations of instant application claim 22 appear to be anticipated or obvious over '505 claims 77 and 82.

For example, instant application claim 22 recites, "...a first semiconductor material comprising a planarized insulating material having a first bonding surface; and a second material having a second bonding surface, wherein the first bonding surface is an etched surface terminated with a nitrogen- containing species, and wherein the first and second bonding surfaces are in direct contact with each other and bonded together with a chemical bond without any intervening adhesive...", which includes the similarly recited limitations in '505 claims 77and 82, *namely*, "...a processed semiconductor device wafer comprising a first surface and a planarized insulating material having a first bonding surface wherein said planarized insulating material is opposite said first surface; a first material having a second surface and a second bonding surface opposite said second surface; wherein said first bonding surface is an etched surface terminated with a species; and wherein said first and second bonding surfaces are in direct contact

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with each other and bonded together with a chemical bond without any adhesive...", and "...wherein said species comprises nitrogen...".

Claims 22-31 are rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over claim 115, of U.S. Patent 8,153,505.

Although the conflicting claims are not identical, it is not patentably distinct from each other because the claims of the current application, U.S. Patent Application 14/957,501, are broader than the claims of the U.S. Patent 8,153,505. In this regard, the limitations of instant application claims 22 and 27, appear to be anticipated or obvious over '505 claim 115.

For example, instant application claim 22 recites, "...a first semiconductor material comprising a planarized insulating material having a first bonding surface; and a second material having a second bonding surface, wherein the first bonding surface is an etched surface terminated with a nitrogen- containing species, and wherein the first and second bonding surfaces are in direct contact with each other and bonded together with a chemical bond without any intervening adhesive...", and "...wherein said planarized insulating material comprises silicon oxide...", which includes the similarly recited limitations in '505 claim 115, *namely*, "...a processed semiconductor device wafer comprising a first surface having a thickness, a planarized insulating material of deposited silicon oxide having a first bonding surface, wherein said planarized insulating material is opposite said first surface, and a device layer between said first surface and said planarized insulating material; a first material comprising silicon having a second

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surface and a second bonding surface opposite said second surface; wherein said first bonding surface is an etched surface terminated with nitrogen; and wherein said first and second bonding surfaces are in direct contact with each other and bonded together with a chemical bond without any adhesive...".

Claims 22-31 are rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over claims 192 and 197, of U.S. Patent 8,153,505. Although the conflicting claims are not identical, it is not patentably distinct from each other because the claims of the current application, U.S. Patent Application 14/957,501, are broader than the claims of the U.S. Patent 8,153,505. In this regard, the limitations of instant application claim 22 appear to be anticipated or obvious over '505 claims 192 and 197.

For example, instant application claim 22 recites, "...a first semiconductor material comprising a planarized insulating material having a first bonding surface; and a second material having a second bonding surface, wherein the first bonding surface is an etched surface terminated with a nitrogen- containing species, and wherein the first and second bonding surfaces are in direct contact with each other and bonded together with a chemical bond without any intervening adhesive...", which includes the similarly recited limitations in '505 claims 192 and 197, *namely*, "...a first silicon material comprising a first surface and a planarized insulating material having a first bonding surface wherein said planarized insulating material is opposite said first surface; a second material having a second surface and a second bonding surface opposite said

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second surface; wherein said first bonding surface is an etched surface terminated with a species; and wherein said first and second bonding surfaces are in direct contact with each other and bonded together with a chemical bond without any adhesive...", and "...wherein said species comprises nitrogen...".

Claims 22-31 are rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over claim 230, of U.S. Patent 8,153,505.

Although the conflicting claims are not identical, it is not patentably distinct from each other because the claims of the current application, U.S. Patent Application 14/957,501, are broader than the claims of the U.S. Patent 8,153,505. In this regard, the limitations of instant application claims 22 and 27, appear to be anticipated or obvious over '505 claim 230.

For example, instant application claim 22 recites, "...a first semiconductor material comprising a planarized insulating material having a first bonding surface; and a second material having a second bonding surface, wherein the first bonding surface is an etched surface terminated with a nitrogen- containing species, and wherein the first and second bonding surfaces are in direct contact with each other and bonded together with a chemical bond without any intervening adhesive...", and "...wherein said planarized insulating material comprises silicon oxide...", which includes the similarly recited limitations in '505 claim 230, *namely*, "...a first silicon material comprising a first surface having a thickness, a planarized insulating material of deposited silicon oxide having a first bonding surface, wherein said planarized insulating material is opposite

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said first surface, and a device layer between said first surface and said planarized insulating material; a second material comprising silicon having a second surface and a second bonding surface opposite said second surface; wherein said first bonding surface is an etched surface terminated with nitrogen; wherein said first and second bonding surfaces are in direct contact with each other and bonded together with a chemical bond without any adhesive...".

Claims 2-21, and 32-34, are rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over claims 1 and 9, of U.S. Patent 9,082,627. Although the conflicting claims are not identical, it is not patentably distinct from each other because the claims of the current application, U.S. Patent Application 14/957,501, are broader than the claims of the U.S. Patent 9,082,627. In this regard, the limitations of instant application claim 2 appear to be anticipated or obvious over '627 claims 1 and 9.

For example, instant application claim 1 recites, "...forming a planarized insulating material on a semiconductor wafer; etching the planarized insulating material in a reaction space; terminating the planarized insulating material with a nitrogen-containing species by at least one of: a nitrogen-containing plasma during the etching, and a nitrogen-containing solution after the etching; bringing a first material into direct contact with the terminated and planarized insulating material outside the reaction space; and forming a chemical bond between the terminated and planarized insulating material and the first material...", which includes the similarly recited limitations in '627

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claims 1 and 9, *namely*, "...forming a planarized insulating material on a processed semiconductor wafer; exposing said planarized insulating material to a plasma RIE process in a plasma chamber; terminating said planarized insulating material with a species; bringing into direct contact said terminated and planarized insulating material and a first material outside of said plasma chamber; and forming a chemical bond between said terminated and planarized insulating material and said first material...", and "...exposing said insulating material to a plasma RIE process in said plasma chamber using nitrogen gas...".

Claims 2-21, and 32-34, are rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over claims 1 and 9, of U.S. Patent 9,082,627. Although the conflicting claims are not identical, it is not patentably distinct from each other because the claims of the current application, U.S. Patent Application 14/957,501, are broader than the claims of the U.S. Patent 9,082,627. In this regard, the limitations of instant application claim 32 appear to be anticipated or obvious over '627 claims 1 and 9.

For example, instant application claim 32 recites, "...forming a planarized insulating material on a semiconductor wafer; etching the planarized insulating material; terminating the planarized insulating material with a nitrogen-containing species by at least one of: a nitrogen-containing plasma during the etching, and a nitrogen-containing solution after the etching; bringing a first material into direct contact with the terminated and planarized insulating material; and forming a chemical bond between the

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terminated and planarized insulating material and the first material...", which includes the similarly recited limitations in '627 claims 1 and 9, *namely*, "...forming a planarized insulating material on a processed semiconductor wafer; exposing said planarized insulating material to a plasma RIE process in a plasma chamber; terminating said planarized insulating material with a species; bringing into direct contact said terminated and planarized insulating material and a first material outside of said plasma chamber; and forming a chemical bond between said terminated and planarized insulating material and said first material...", and "...exposing said insulating material to a plasma RIE process in said plasma chamber using nitrogen gas...".

Claims 2-21, and 32-34, are rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over claim 1, of U.S. Patent 9,082,627. Although the conflicting claims are not identical, it is not patentably distinct from each other because the claims of the current application, U.S. Patent Application 14/957,501, are broader than the claims of the U.S. Patent 9,082,627. In this regard, the limitations of instant application claim 33 appear to be anticipated or obvious over '627 claim 1.

For example, instant application claim 33 recites, "...forming a planarized insulating material on a processed semiconductor wafer; exposing the planarized insulating material t a plasma process in a plasma chamber using a nitrogen-containing gas; terminating the planarized insulating material with a species; bringing into direct contact the terminated and planarized insulating material and a first material outside of

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said plasma chamber; and forming a chemical bond between the terminated and planarized insulating material and the first material...", which includes the similarly recited limitations in '627 claim 1, *namely*, "...forming a planarized insulating material on a processed semiconductor wafer; exposing said planarized insulating material to a plasma RIE process in a plasma chamber; terminating said planarized insulating material with a species; bringing into direct contact said terminated and planarized insulating material and a first material outside of said plasma chamber; and forming a chemical bond between said terminated and planarized insulating material and said first material...".

Claims 2-21, and 32-34, are rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over claims 1, 2, and 11, of U.S. Patent 8,153,505. Although the conflicting claims are not identical, it is not patentably distinct from each other because the claims of the current application, U.S. Patent Application 14/957,501, are broader than the claims of the U.S. Patent 9,082,627. In this regard, the limitations of instant application claim 32 appear to be anticipated or obvious over '627 claims 1, 2, and 11.

For example, instant application claim 32 recites, "...forming a planarized insulating material on a semiconductor wafer; etching the planarized insulating material; terminating the planarized insulating material with a nitrogen-containing species by at least one of: a nitrogen-containing plasma during the etching, and a nitrogen-containing solution after the etching; bringing a first material into direct contact with the terminated

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and planarized insulating material; and forming a chemical bond between the terminated and planarized insulating material and the first material...", which includes the similarly recited limitations in '627 claims 1 and 9, *namely*, "...forming a planarized insulating material on a processed semiconductor wafer; exposing said planarized insulating material to a plasma RIE process; terminating said planarized insulating material with a species; after said exposing and terminating steps, bringing into direct contact said terminated planarized insulating material and a first material; and forming a chemical bond between said terminated planarized insulating material and said first material...", "...wherein said chemical bond is formed without any adhesive...", and "...wherein said plasma RIE process comprises a nitrogen gas...".

Claims 2-21, and 32-34, are rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over claims 39, 40, and 48, of U.S. Patent 8,153,505. Although the conflicting claims are not identical, it is not patentably distinct from each other because the claims of the current application, U.S. Patent Application 14/957,501, are broader than the claims of the U.S. Patent 8,153,505. In this regard, the limitations of instant application claim 32 appear to be anticipated or obvious over '505 claims 39, 40, and 48.

For example, instant application claim 32 recites, "...forming a planarized insulating material on a semiconductor wafer; etching the planarized insulating material; terminating the planarized insulating material with a nitrogen-containing species by at least one of: a nitrogen-containing plasma during the etching, and a nitrogen-containing

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solution after the etching; bringing a first material into direct contact with the terminated and planarized insulating material; and forming a chemical bond between the terminated and planarized insulating material and the first material...", which includes the similarly recited limitations in '505 claims 39, 40, and 48, *namely*, "...forming a planarized insulating material on a processed semiconductor wafer; exposing said planarized insulating material to a plasma RIE process; terminating said planarized insulating material with a first species; terminating a first material with a second species; bringing into direct contact said terminated planarized insulating material with said terminated first material; and forming a chemical bond between said terminated planarized insulating material and said terminated first material...", "...wherein said chemical bond is formed without any adhesive...", and "...wherein said plasma RIE process contains a nitrogen gas...".

Claims 2-21, and 32-34, are rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over claims 116, 117, and 126, of U.S. Patent 8,153,505. Although the conflicting claims are not identical, it is not patentably distinct from each other because the claims of the current application, U.S. Patent Application 14/957,501, are broader than the claims of the U.S. Patent 8,153,505. In this regard, the limitations of instant application claim 32 appear to be anticipated or obvious over '505 claims 116, 117, and 126.

For example, instant application claim 32 recites, "...forming a planarized insulating material on a semiconductor wafer; etching the planarized insulating material;

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terminating the planarized insulating material with a nitrogen-containing species by at least one of: a nitrogen-containing plasma during the etching, and a nitrogen-containing solution after the etching; bringing a first material into direct contact with the terminated and planarized insulating material; and forming a chemical bond between the terminated and planarized insulating material and the first material...", which includes the similarly recited limitations in '505 claims 116, 117, and 126, namely, "...forming a planarized insulating material on a first silicon material; exposing said planarized insulating material to a plasma RIE process; terminating said planarized insulating material with a species; after exposing and terminating steps, bringing into direct contact said terminated planarized insulating material and a second material; and forming a chemical bond between said terminated planarized insulating material and said second material...", "...wherein said chemical bond is formed without any adhesive...", and "...wherein said plasma RIE process contains a nitrogen gas...".

Claims 2-21, and 32-34, are rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over claims 154, 155, and 163, of U.S. Patent 8,153,505. Although the conflicting claims are not identical, it is not patentably distinct from each other because the claims of the current application, U.S. Patent Application 14/957,501, are broader than the claims of the U.S. Patent 8,153,505. In this regard, the limitations of instant application claim 32 appear to be anticipated or obvious over '505 claims 154, 155, and 163.

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For example, instant application claim 32 recites, "...forming a planarized insulating material on a semiconductor wafer; etching the planarized insulating material; terminating the planarized insulating material with a nitrogen-containing species by at least one of: a nitrogen-containing plasma during the etching, and a nitrogen-containing solution after the etching; bringing a first material into direct contact with the terminated and planarized insulating material; and forming a chemical bond between the terminated and planarized insulating material and the first material...", which includes the similarly recited limitations in '505 claims 154, 155, and 163, namely, "...forming a planarized insulating material on a first silicon material; exposing said planarized insulating material to a plasma RIE process; terminating said planarized insulating material with a first species; terminating a second material with a second species; bringing into direct contact said terminated planarized insulating material with said terminated second material; and forming a chemical bond between said terminated planarized insulating material and said terminated second material...", "...wherein said chemical bond is formed without any adhesive...", and "...wherein said plasma RIE process contains a nitrogen gas...".

Claims 2-21, and 32-34, are rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over claims 105, 106, and 107, of U.S. Patent 6,902,987. Although the conflicting claims are not identical, it is not patentably distinct from each other because the claims of the current application, U.S. Patent Application 14/957,501, are broader than the claims of the U.S. Patent

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6,902,987. In this regard, the limitations of instant application claim 32 appear to be anticipated or obvious over '987 claims 105, 106, and 107.

For example, instant application claim 32 recites, "...forming a planarized insulating material on a semiconductor wafer; etching the planarized insulating material; terminating the planarized insulating material with a nitrogen-containing species by at least one of: a nitrogen-containing plasma during the etching, and a nitrogen-containing solution after the etching; bringing a first material into direct contact with the terminated and planarized insulating material; and forming a chemical bond between the terminated and planarized insulating material and the first material...", which includes the similarly recited limitations in '5987claims 105, 106, and 107, namely, "...depositing first and second bonding layers on respective first and second workpieces; planarizing said first and second bonding layers; etching said first and second bonding layers after planarizing to activate respective first and second surfaces of said first and second bonding layers; terminating said first and second surfaces with a terminating chemical species after etching; bringing into direct contact said first and second surfaces after said etching step; and forming a chemical bond between said first and second workpieces at room temperature...", "...depositing each of said first and second bonding layers comprises depositing a silicon oxide layer; planarizing comprises polishing said first and second bonding layers; etching comprises exposing said first and second bonding layers to a plasma; and terminating comprises immersing said first and second bonding surfaces in an ammonia-based solution...", and "...wherein said first and second workpieces each comprise a semiconductor wafer...".

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Conclusion

When responding to this office action, applicants are advised to provide the

examiner with the line numbers and page numbers in the application and/or references

cited to assist the examiner in locating appropriate paragraphs.

Any inquiry concerning this communication or earlier communications from the

examiner should be directed to Joannie García, whose telephone number is (571) 272-

1861. The examiner can normally be reached on Monday through Friday.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's

supervisor, Mohammad K. Islam, can be reached on (571) 270-5878. The fax phone

number for the organization where this application or proceeding is assigned is 571-

273-8300.

Information regarding the status of an application may be obtained from the

Patent Application Information Retrieval (PAIR) system. Status information for

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/JOANNIE A GARCIA/

Examiner, Art Unit 2895

/KYOUNG LEE/

Primary Examiner, Art Unit 2895

Application/Control No. Applicant(s)/Patent Under Reexamination 14/957,501 TONG ET AL. Notice of References Cited Examiner Art Unit Page 1 of 1 JOANNIE A. GARCIA 2895 **U.S. PATENT DOCUMENTS** Document Number Date **CPC Classification US Classification** Name Country Code-Number-Kind Code MM-YYYY * 05-2002 US-2002/0055208 A1 Ohtani, Hisashi H01L21/2022 438/151 Α В US-US-С US-D US-Ε US-F US-G Н US-US-US-J US-Κ US-L US-Μ

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U.S. Patent and Trademark Office PTO-892 (Rev. 01-2001)

Notice of References Cited

Part of Paper No. 01252016

	Application No.	14/957,501
INFORMATION DISCLOSURE	Filing Date	December 2, 2015
STATEMENT BY APPLICANT	First Named Inventor	Tong et al.
STATEMENT BY AFFEIGANT	Art Unit	2895
(Multiple sheets used when necessary)	Examiner	Unknown
SHEET 1 OF 5	Attorney Docket No.	TSSRA.012C1

	U.S. PATENT DOCUMENTS						
Examiner Initials	Cite No.	Document Number Number - Kind Code (if known) Example: 1,234,567 B1	Publication Date MM-DD-YYYY	Name	Pages, Columns, Lines Where Relevant Passages or Relevant Figures Appear		
	1	4,700,466	10-20-1987	Nakagawa et al.			
	2	4,983,251	01-08-1991	Haisma et al.			
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Examiner Initials	Cite No.	Foreign Patent Document Country Code-Number-Kind Code Example: JP 1234567 A1	Publication Date MM-DD-YYYY	Name	Pages, Columns, Lines Where Relevant Passages or Relevant Figures Appear	T ¹		
	5	CA 2 399 282 A1	08-23-2001	Ziptronix, Inc.				
	6	JP H08-195334	07-30-1996	Canon Inc.		Т		
	7	TW 389965B	05-11-2000	San Fu Chemical Co. Ltd.		Т		

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SHEET 2 OF 5	Attorney Docket No.	TSSRA.012C1

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SHEET 3 OF 5	Attorney Docket No.	TSSRA.012C1

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PTO/SB/08 Equivalent

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EAST Search History

EAST Search History (Prior Art)

Ref #	Hits	Search Query	DBs	Default Operator	Plurals	Time Stamp
L1	472066	(bond\$).clm.	US- PGPUB; USPAT; USOCR	OR	OFF	2016/01/25 18:29
L2	58187	(anneal\$).clm.	US- PGPUB; USPAT; USOCR	OR	OFF	2016/01/25 18:29
L3	865929	(heat or heats or heated or heating).clm.	US- PGPUB; USPAT; USOCR	• 5	OFF	2016/01/25 18:29
L4	140	L1 with L2 with L3	US- PGPUB; USPAT; USOCR	OR	OFF	2016/01/25 18:29
L5	22	L4 AND ((H01L21/324 OR H01L21/2022 OR H01L21/26506 OR H01L21/187 OR H01L21/2007 OR H01L21/76251 OR H01L23/38 OR H01L21/02032 OR H01L21/3065 OR H01L21/3226 OR H01L21/67092).CPC.)	US- PGPUB; USPAT; USOCR		OFF	2016/01/25 18:29
L6	19956	((257/459) or (257/676) or (257/684) or (257/701) or (257/702) or (257/e21.088) or (257/e21.122) or (257/e21.215) or (257/e21.218) or (257/e21.219) or (257/e21.224) or (257/e21.224) or (257/e21.243) or (257/e21.245) or (257/e21.246) or (257/e21.249) or (257/e21.33) or (257/e21.567) or (257/e23.02) or (257/e23.04)).CCLS.	US- PGPUB; USPAT; USOCR		OFF	2016/01/25 18:45
L7	5551	((438/405) or (438/455) or (438/459) or (438/974) or (148/DIG12)).CCLS.	US- PGPUB; USPAT; USOCR		OFF	2016/01/25 18:51
L8	24326	6 or 7	US- PGPUB; USPAT; USOCR		OFF	2016/01/25 18:51
L9	1406	(bond or bonds or bonded or bonding) with (etch or etches or etched or etching) with ("25" or "room temperature" or "room-temperature" or "room-T")	US- PGPUB; USPAT; USOCR		OFF	2016/01/25 18:52
L10	28	(bond or bonds or bonded or bonding) with (etch or etches or etched or etching) with (("25" or "room temperature" or "room-temperature" or "room T" or "room-T") and ("chemical bond" or "chemical-bond"))	US- PGPUB; USPAT; USOCR	OR	ON	2016/01/25 18:53
L11	4	(("4700466") or ("4983251") or	US-	OR	OFF	2016/01/25

		("5427638") or ("5451547")).PN.	PGPUB; USPAT; USOCR			18:58
L12	1	("20150303263").PN.	US- PGPUB; USPAT; USOCR	OR	OFF	2016/01/25 19:30
L13	86	"planarized insulating material"	US- PGPUB; USPAT; USOCR	OR	ON	2016/01/25 19:30
L14	1	L12 and L13	US- PGPUB; USPAT; USOCR	OR	ON	2016/01/25 19:30
S1	1	("20140203407").PN.	US- PGPUB; USPAT; USOCR	OR	OFF	2014/10/17 09:41
S2	1	("8153505").PN.	US- PGPUB; USPAT; USOCR	OR	OFF	2014/10/17 09:42
S3	1349	(bond or bonds or bonded or bonding) with (etch or etches or etched or etching) with ("25" or "room temperature" or "room- temperature" or "room T" or "room-T")	US- PGPUB; USPAT; USOCR	OR	OFF	2014/10/17 09:47
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S5	0	(bond or bonds or bonded or bonding) with (etch or etches or etched or etching) with (("25" or "room temperature" or "room-temperature" or "room T" or "room-T") same ("chemical bond" or "chemicalbond"))	US- PGPUB; USPAT; USOCR		OFF	2014/10/17 09:50
S6	5	(bond or bonds or bonded or bonding) with (etch or etches or etched or etching) with (("25" or "room temperature" or "room- temperature" or "room T" or "room-T") and ("chemical bond" or "chemical-bond"))	PGPUB; USPAT;	OR	OFF	2014/10/17 09:51
S7	0	((bond or bonds or bonded or bonding) with (etch or etches or etched or etching) with (("25" or "room temperature" or "room-temperature" or "room T" or "room- T") and ("chemical bond" or "chemical- bond"))).clm.	US- PGPUB; USPAT; USOCR	OR	OFF	2014/10/17 09:51
S8	436228	(bond\$).clm.	US- PGPUB; USPAT; USOCR	OR	OFF	2014/10/17 09:53
S9	53191	(anneal\$).clm.	US- PGPUB; USPAT; USOCR	OR	OFF	2014/10/17 09:53
S10	802974	(heat or heats or heated or heating).clm.	US-	OR	OFF	2014/10/17

			PGPUB; USPAT; USOCR			09:54
S11	134	S8 with S9 with S10	US- PGPUB; USPAT; USOCR	OR	OFF	2014/10/17 09:54
S12	19293	((257/459) or (257/676) or (257/684) or (257/701) or (257/702) or (257/e21.088) or (257/e21.122) or (257/e21.215) or (257/e21.218) or (257/e21.219) or (257/e21.224) or (257/e21.224) or (257/e21.243) or (257/e21.245) or (257/e21.246) or (257/e21.246) or (257/e21.33) or (257/e21.567) or (257/e23.02) or (257/e23.04)).CCLS.	US- PGPUB; USPAT; USOCR	OR	OFF	2014/10/17 10:07
S13	6	S11 and S12	US- PGPUB; USPAT; USOCR	OR	OFF	2014/10/17 10:09
S14	18	S11 AND ((H01L21/324 OR H01L21/2022 OR H01L21/26506 OR H01L21/187 OR H01L21/2007 OR H01L21/76251 OR H01L23/38 OR H01L21/02032 OR H01L21/3065 OR H01L21/3226 OR H01L21/67092).CPC.)	US- PGPUB; USPAT; USOCR	OR	OFF	2014/10/17 10:50
S15	1386	(bond or bonds or bonded or bonding) with (etch or etches or etched or etching) with ("25" or "room temperature" or "room- temperature" or "room T" or "room-T")	US- PGPUB; USPAT; USOCR	OR	OFF	2015/07/22 15:54
S16	0	(bond or bonds or bonded or bonding) with (etch or etches or etched or etching) with (("25" or "room temperature" or "room- temperature" or "room T" or "room-T") with ("chemical bond" or "chemical- bond"))	US- PGPUB; USPAT; USOCR	OR	OFF	2015/07/22 15:54
S17	0	(bond or bonds or bonded or bonding) with (etch or etches or etched or etching) with (("25" or "room temperature" or "room-temperature" or "room T" or "room-T") same ("chemical bond" or "chemicalbond"))	US- PGPUB; USPAT; USOCR	OR	OFF	2015/07/22 15:54
S18	5	(bond or bonds or bonded or bonding) with (etch or etches or etched or etching) with (("25" or "room temperature" or "room- temperature" or "room T" or "room-T") and ("chemical bond" or "chemical-bond"))	PGPUB; USPAT;	OR	OFF	2015/07/22 15:54
S19	0	((bond or bonds or bonded or bonding) with (etch or etches or etched or etching) with (("25" or "room temperature" or "room-temperature" or "room T" or "room- T") and ("chemical bond" or "chemical- bond"))).clm.	US- PGPUB; USPAT; USOCR	OR	OFF	2015/07/22 15:54
S20	457036	(bond\$).clm.	US- PGPUB; USPAT; USOCR	OR	OFF	2015/07/22 15:54
S21	56069	(anneal\$).clm.	US- PGPUB; USPAT;	OR	OFF	2015/07/22 15:54

			USOCR		***************************************	
S22	839493	(heat or heats or heated or heating).clm.	US- PGPUB; USPAT; USOCR	2	OFF	2015/07/22 15:54
S23	138	오0 with 오1 with 오2	US- PGPUB; USPAT; USOCR	OR	OFF	2015/07/22 15:55
S24	19807	((257/459) or (257/676) or (257/684) or (257/701) or (257/702) or (257/e21.088) or (257/e21.122) or (257/e21.215) or (257/e21.218) or (257/e21.219) or (257/e21.224) or (257/e21.243) or (257/e21.245) or (257/e21.246) or (257/e21.249) or (257/e21.33) or (257/e21.567) or (257/e23.02) or (257/e23.04)).CCLS.	US- PGPUB; USPAT; USOCR	,	OFF	2015/07/22 15:55
S25	6	S23 and S24	US- PGPUB; USPAT; USOCR	OR	OFF	2015/07/22 15:55
S26	20	\$23 AND ((H01L21/324 OR H01L21/2022 OR H01L21/26506 OR H01L21/187 OR H01L21/2007 OR H01L21/76251 OR H01L23/38 OR H01L21/02032 OR H01L21/3065 OR H01L21/3226 OR H01L21/67092).CPC.)	US- PGPUB; USPAT; USOCR	OR	OFF	2015/07/22 15:55
S27	465993	(bond\$).clm.	US- PGPUB; USPAT; USOCR	OR	OFF	2015/11/10 16:40
S28	57339	(anneal\$).clm.	US- PGPUB; USPAT; USOCR	OR	OFF	2015/11/10 16:40
S29	855382	(heat or heats or heated or heating).clm.	US- PGPUB; USPAT; USOCR	OR	OFF	2015/11/10 16:40
S30	139	S27 with S28 with S29	US- PGPUB; USPAT; USOCR	OR	OFF	2015/11/10 16:40
S31	21	\$30 AND ((H01L21/324 OR H01L21/2022 OR H01L21/26506 OR H01L21/187 OR H01L21/2007 OR H01L21/76251 OR H01L23/38 OR H01L21/02032 OR H01L21/3065 OR H01L21/3226 OR H01L21/67092).CPC.)	US- PGPUB; USPAT; USOCR	OR	OFF	2015/11/10 16:40
S32	1	("20150303263").PN.	US- PGPUB; USPAT; USOCR	OR	OFF	2015/11/10 16:41
S33	1	("5451547").PN.	US- PGPUB; USPAT; USOCR	OR	OFF	2015/11/10 16:41
S34	1286659	adhesive	US-	OR	ON	2015/11/10

			PGPUB; USPAT; USOCR			16:41
S35	0	S33 and S34	US- PGPUB; USPAT; USOCR	OR	ON	2015/11/10 16:41
S36	2043	terminated with species	US- PGPUB; USPAT; USOCR	OR	ON	2015/11/10 16:45
S37	1	S32 and S36	US- PGPUB; USPAT; USOCR	OR	ON	2015/11/10 16:45
S38	909961	etch\$	US- PGPUB; USPAT; USOCR	OR	ON	2015/11/10 16:48
S39	1	S33 and S38	US- PGPUB; USPAT; USOCR	OR	ON	2015/11/10 16:49
S40	5	(bond or bonds or bonded or bonding) with (etch or etches or etched or etching) with (("25" or "room temperature" or "room- temperature" or "room T" or "room-T") and ("chemical bond" or "chemical-bond"))	PGPUB; USPAT;	OR	OFF	2015/11/10 16:58
S41	28	(bond or bonds or bonded or bonding) with (etch or etches or etched or etching) with (("25" or "room temperature" or "room-temperature" or "room T" or "room-T") and ("chemical bond" or "chemical-bond"))	PGPUB; USPAT;	OR	ON	2015/11/10 16:58
S42	0	S33 and S41	US- PGPUB; USPAT; USOCR	OR	ON	2015/11/10 16:58
S43	48227	(bond or bonds or bonded or bonding) with (etch or etches or etched or etching)	US- PGPUB; USPAT; USOCR	OR	ON	2015/11/10 16:58
S44	0	S33 and S43	US- PGPUB; USPAT; USOCR	OR	ON	2015/11/10 16:59
S45	2042101	bond or bonds or bonded or bonding	US- PGPUB; USPAT; USOCR	OR	ON	2015/11/10 16:59
S46	1	S33 and S45	US- PGPUB; USPAT; USOCR	OR	ON	2015/11/10 16:59
S47	567781	CMP or polish\$ or planarize or planarizes or planarized or planarizing or planarization	US- PGPUB; USPAT; USOCR	OR	ON	2015/11/10 17:04
S48		S33 and S47	US- PGPUB;	OR	ON	2015/11/10 17:04

			USPAT; USOCR	***************************************	
S49	3009473	plasma or gas	US- PGPUB; USPAT; USOCR	ON	2015/11/10 17:09
S50	1	S33 and S49	US- PGPUB; USPAT; USOCR	ON	2015/11/10 17:09
S51	1	S39 and S46 and S48 and S50	US- PGPUB; USPAT; USOCR	ON	2015/11/10 17:10
S52	913511	etch\$ or VSE	US- PGPUB; USPAT; USOCR	ON	2015/11/10 17:49
S53	1	S33 and S52	US- PGPUB; USPAT; USOCR	ON	2015/11/10 17:49
S54	790750	specie	US- PGPUB; USPAT; USOCR	ON	2015/11/10 17:53
S55	0	S33 and S54	US- PGPUB; USPAT; USOCR	ON	2015/11/10 17:54
S56	1	("5451547").PN.	US- PGPUB; USPAT; USOCR	OFF	2015/11/14 10:09
S57	1	("20150303263").PN.	US- PGPUB; USPAT; USOCR	OFF	2015/11/14 10:38
S58	85	"planarized insulating material"	US- PGPUB; USPAT; USOCR	ON	2015/11/14 10:43
S59	1	S57 and S58	US- PGPUB; USPAT; USOCR	ON	2015/11/14 10:43

1/25/2016 7:31:19 PM

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UNITED STATES PATENT AND TRADEMARK OFFICE

UNITED STATES DEPARTMENT OF COMMERCE United States Patent and Trademark Office Address: COMMISSIONER FOR PATENTS P.O. Box 1450 Alexandria, Virginia 22313-1450 www.uspto.gov

BIB DATA SHEET

CONFIRMATION NO. 1033

SERIAL NUM	IRED	FILING or	371(c)		CLASS	GB		LINIT	ΔΤΤΟ	DRNEY DOCKET
14/957,50		12/02/2	E `´		257	un	2895		NO. TSSRA.012C1	
14/001,00	, ,	RUL			201		2000		'	33NA.012G1
	APPLICANTS									
	·	C., Morrisville,	NC;							
Qin-Yi To Gaius Gil	INVENTORS Qin-Yi Tong, Durham, NC; Gaius Gillman Fountain JR., Youngsville, NC; Paul M. Enquist, Cary, NC;									
This appl wh wh wh wh wh wh	** CONTINUING DATA **********************************									
** FOREIGN A	PPLICA	ATIONS *****	******	*****	*					
** IF REQUIRE 12/11/20		REIGN FILING	LICENS	E GRA	ANTED **					
Foreign Priority claims	ditions me		☐ Met at Allowa	ter ance	STATE OR COUNTRY		HEETS WINGS	TOT.		INDEPENDENT CLAIMS
	/JOANNIE Examiner's	A GARCIA/ Signature	JAG Initials		NC		9	33	3	4
ADDRESS	ADDRESS									
Knobbe, Martens, Olson & Bear, LLP 2040 main Street Fourteenth Floor Irvine, CA 92614 UNITED STATES										
TITLE										
METHOD FOR LOW TEMPERATURE BONDING AND BONDED STRUCTURE										
☐ All Fees										
FILING FEE RECEIVED 3200 FEES: Authority has been given in Paper No to charge/credit DEPOSIT ACCOUNT No for following: 1.16 Fees (Filing) 1.17 Fees (Processing Ext. of to the processing Ext. of the processing Ext										
					ing Ext. of time)					
							☐ Other	· 		
☐ Credit										

BIB (Rev. 05/07).

Search Notes Application/Control No. Applicant(s)/Patent Under Reexamination TONG ET AL. Examiner JOANNIE A GARCIA Applicant(s)/Patent Under Reexamination TONG ET AL.

CPC- SEARCHED		
Symbol	Date	Exa mine r
H01L: 21/02032,21/185,21/187,21/2007,21/2022,21/26506,21/30,21/302,21/306,21/3065,21/3226,21/324,21/38,21/46,21/461,21/67092,21/76251,21/76254	01/25/ 2016	JAG

CPC COMBINATION SETS - SEARCHED				
Symbol	Date	Examiner		

US CLASSIFICATION SEARCHED				
Class	Subclass	Date	Examiner	
438	405,455,459,974,FOR103,FOR127,FOR388,FOR395	01/25/2016	JAG	
257	459,676,684,701,702,E21.088,E21.122,E21.215,E21.218 ,E21.219,E21.224,E21.243,E21.245,E21.249,E21.33,E23 .02,E23.04	01/25/2016	JAG	
148	DIG12	01/25/2016	JAG	

SEARCH NOTES		
Search Notes	Date	Examiner
EAST search attached	01/25/2016	JAG

INTERFERENCE SEARCH					
US Class/ CPC Symbol	US Subclass / CPC Group	Date	Examiner		
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Doc Code: DIST.E.FILE Document Description: Electronic To	erminal Disclaimer - Filed		PTO/SB/25 PTO/SB/26 U.S. Patent and Trademark Office Department of Commerce		
Electronic Petition Request	TERMINAL DISCLAIMER TO OBVIATE A PROVISIONAL DOUBLE PATENTING REJECTION OVER A PENDING "REFERENCE" APPLICATION AND TERMINAL DISCLAIMER TO OBVIATE A DOUBLE PATENTING REJECTION OVER A "PRIOR" PATENT				
Application Number	14957501				
Filing Date	02-Dec-2015				
First Named Inventor	Qin-Yi Tong				
Attorney Docket Number	TSSRA.012C1				
Title of Invention	METHOD FOR LOW TEMPERATURE BONDING AND BONDED STRUCTURE				
Filing of terminal disclaimer does Office Action	not obviate requirement for res	ponse under 37	CFR 1.111 to outstanding		
This electronic Terminal Disclaim	er is not being used for a Joint Re	esearch Agreem	ent.		
Owner	P	ercent Interest			
ZIPTRONIX, INC.	1	00 %			

The owner(s) of percent interest listed above in the instant application hereby disclaims, except as provided below, the terminal part of the statutory term of any patent granted on the instant application which would extend beyond the expiration date of the full statutory term of any patent granted on pending reference Application Number(s)

14754111 filed on 06/29/2015

as the term of any patent granted on said reference application may be shortened by any terminal disclaimer filed prior to the grant of any patent on the pending reference application. The owner hereby agrees that any patent so granted on the instant application shall be enforceable only for and during such period that it and any patent granted on the reference application are commonly owned. This agreement runs with any patent granted on the instant application and is binding upon the grantee, its successors or assigns.

In making the above disclaimer, the owner does not disclaim the terminal part of any patent granted on the instant application that would extend to the expiration date of the full statutory term of any patent granted on said reference application, "as the term of any patent granted on said reference application may be shortened by any terminal disclaimer filed prior to the grant of any patent on the pending reference application," in the event that any such patent granted on the pending reference application: expires for failure to pay a maintenance fee, is held unenforceable, is found invalid by a court of competent jurisdiction, is statutorily disclaimed in whole or terminally disclaimed under 37 CFR 1.321, has all claims canceled by a reexamination certificate, is reissued, or is in any manner terminated prior to the expiration of its full statutory term as shortened by any terminal disclaimer filed prior to its grant.

The owner(s) with percent interest listed above in the instant application hereby disclaims, except as provided below, the terminal part of the statutory term of any patent granted on the instant application which would extend beyond the expiration

Page र्मान् भारिकापारा term of prior patent number(s)

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granted on the instant application :	resently shortened by any terminal disclaimer. The owner hereby agrees that any patent so shall be enforceable only for and during such period that it and the prior patent are commonly ny patent granted on the instant application and is binding upon the grantee, its successors
application that would extend to the is presently shortened by any terming expires for failure to pay a maintener is held unenforceable; - is found invalid by a court of comparts of the invalid by a court of comparts of the invalid by a court of comparts of the invalid by a court of the invalid by a court of the invalid by a court of comparts of the invalid by a court of the inva	petent jurisdiction; or terminally disclaimed under 37 CFR 1.321;
- is reissued; or	mination certificate,
- is in any manner terminated prior	to the expiration of its full statutory term as presently shortened by any terminal disclaimer.
Terminal disclaimer fee unde	r 37 CFR 1.20(d) is included with Electronic Terminal Disclaimer request.
I/ 1	7 CFR 1.4(d)(4), that the terminal disclaimer fee under 37 CFR 1.20(d) claimer has already been paid in the above-identified application.
Applicants claims the following fee	status:
Small Entity	
Micro Entity	
Regular Undiscounted	
belief are believed to be true; and for the like so made are punishable by	made herein of my own knowledge are true and that all statements made on information and urther that these statements were made with the knowledge that willful false statements and fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and ay jeopardize the validity of the application or any patent issued thereon.
THIS PORTION MUST BE COMPLET	ED BY THE SIGNATORY OR SIGNATORIES
I certify, in accordance with 37 CFI	R 1.4(d)(4) that I am:
An attorney or agent registere this application	ed to practice before the Patent and Trademark Office who is of record in
Registration Number 6828	37
A sole inventor	
A joint inventor; I certify that I power of attorney in the appl	am authorized to sign this submission on behalf of all of the inventors as evidenced by the ication
A joint inventor; all of whom a	are signing this request
Signature	/Paul Stellman/
age 216 of 331	SAMSUNG EXHIBIT 100

Tau semian	Name Pa	aul Stellman
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*Statement under 37 CFR 3.73(b) is required if terminal disclaimer is signed by the assignee (owner). Form PTO/SB/96 may be used for making this certification. See MPEP § 324.

Electronic Patent Application Fee Transmittal					
Application Number:	149	957501			
Filing Date:	02-Dec-2015				
Title of Invention:	ME	THOD FOR LOW TE	MPERATURE BO	ONDING AND BONE	DED STRUCTURE
First Named Inventor/Applicant Name:	Qin-Yi Tong				
Filer:	Paul Steven Stellman/Anthony Bonilla				
Attorney Docket Number:	TSSRA.012C1				
Filed as Large Entity					
Filing Fees for Utility under 35 USC 111(a)					
Description		Fee Code	Quantity	Amount	Sub-Total in USD(\$)
Basic Filing:					
Statutory or Terminal Disclaimer		1814	1	160	160
Pages:				•	
Claims:					
Miscellaneous-Filing:					
Petition:					
Patent-Appeals-and-Interference:					
Post-Allowance-and-Post-Issuance:					

Description	Fee Code	Quantity	Amount	Sub-Total in USD(\$)
Extension-of-Time:				
Miscellaneous:				
	Tot	al in USD	(\$)	160

Doc Code: DISQ.E.FILE Document Description: Electronic Terminal Disclaimer – Approved
Application No.: 14957501
Filing Date: 02-Dec-2015
Applicant/Patent under Reexamination: Tong et al.
Electronic Terminal Disclaimer filed on February 8, 2016
This patent is subject to a terminal disclaimer
DISAPPROVED
Approved/Disapproved by: Electronic Terminal Disclaimer automatically approved by EFS-Web
U.S. Patent and Trademark Office

Electronic Acknowledgement Receipt			
EFS ID:	24854569		
Application Number:	14957501		
International Application Number:			
Confirmation Number:	1033		
Title of Invention:	METHOD FOR LOW TEMPERATURE BONDING AND BONDED STRUCTURE		
First Named Inventor/Applicant Name:	Qin-Yi Tong		
Customer Number:	135980		
Filer:	Paul Steven Stellman/Anthony Bonilla		
Filer Authorized By:	Paul Steven Stellman		
Attorney Docket Number:	TSSRA.012C1		
Receipt Date:	08-FEB-2016		
Filing Date:	02-DEC-2015		
Time Stamp:	18:23:16		
Application Type:	Utility under 35 USC 111(a)		

Payment information:

Submitted with Payment	yes
Payment Type	Credit Card
Payment was successfully received in RAM	\$160
RAM confirmation Number	5749
Deposit Account	111410
Authorized User	KNOBBE MARTENS OLSON AND BEAR

The Director of the USPTO is hereby authorized to charge indicated fees and credit any overpayment as follows:

Charge any Additional Fees required under 37 CFR 1.16 (National application filing, search, and examination fees)

File Listing	j:				
Document Number	Document Description	File Name	File Size(Bytes)/ Message Digest	Multi Part /.zip	Pages (if appl.)
1	Flantus via Tourninal Disalaine ou Filad	e Terminal - Disclaimer.pdf	37207	no	3
1	Electronic Terminal Disclaimer-Filed		ab936ebc4ecd29254e40fcd88d5aa6c0a24 3df43		
Warnings:			1		
Information:					
2	Fee Worksheet (SB06)	fee-info.pdf	30677	no no	2
-	ree worldness (5500)		346f24103b946738312373fd73cf1a1ade06 f6d5	1.0	-
Warnings:	<u>.</u>		. ,	•	
Information:					
		Total Files Size (in bytes)	6	7884	

This Acknowledgement Receipt evidences receipt on the noted date by the USPTO of the indicated documents, characterized by the applicant, and including page counts, where applicable. It serves as evidence of receipt similar to a Post Card, as described in MPEP 503.

New Applications Under 35 U.S.C. 111

If a new application is being filed and the application includes the necessary components for a filing date (see 37 CFR 1.53(b)-(d) and MPEP 506), a Filing Receipt (37 CFR 1.54) will be issued in due course and the date shown on this Acknowledgement Receipt will establish the filing date of the application.

National Stage of an International Application under 35 U.S.C. 371

If a timely submission to enter the national stage of an international application is compliant with the conditions of 35 U.S.C. 371 and other applicable requirements a Form PCT/DO/EO/903 indicating acceptance of the application as a national stage submission under 35 U.S.C. 371 will be issued in addition to the Filing Receipt, in due course.

New International Application Filed with the USPTO as a Receiving Office

If a new international application is being filed and the international application includes the necessary components for an international filing date (see PCT Article 11 and MPEP 1810), a Notification of the International Application Number and of the International Filing Date (Form PCT/RO/105) will be issued in due course, subject to prescriptions concerning national security, and the date shown on this Acknowledgement Receipt will establish the international filing date of the application.

TSSRA.012C1 PATENT

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Inventor : Tong et al.

App. No. : 14/957,501

Filed : December 2, 2015

For : METHOD FOR LOW

TEMPERATURE BONDING AND

BONDED STRUCTURE

Examiner : Joannie A. Garcia

Art Unit : 2895

Conf. No. : 1033

RESPONSE TO NON-FINAL OFFICE ACTION

Mail Stop Amendment

Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450

Dear Sir:

In response to the Office Action dated February 3, 2016, Applicant submits the following remarks.

A Listing of Claims begins on page 2 of this paper.

Remarks/Arguments begin on page 6 of this paper.

Filing Date: December 2, 2015

LISTING OF THE CLAIMS

1. (Canceled)

2. (Previously Presented) A bonding method, comprising:

forming a planarized insulating material on a semiconductor wafer;

etching the planarized insulating material in a reaction space;

terminating the planarized insulating material with a nitrogen-containing species

by at least one of:

a nitrogen-containing plasma during the etching, and

a nitrogen-containing solution after the etching;

bringing a first material into direct contact with the terminated and planarized

insulating material outside the reaction space; and

forming a chemical bond between the terminated and planarized insulating

material and the first material.

3. (Previously Presented) The method of Claim 2, wherein the etching the

planarized insulating material is conducted in a vacuum chamber, the reaction space comprising

the vacuum chamber.

4. (Previously Presented) The method of Claim 2, wherein the etching the

planarized insulating material is conducted in a plasma chamber, the reaction space comprising

the plasma chamber.

5. (Previously Presented) The method of Claim 2, wherein the etching and the

terminating comprise exposing the planarized insulating material to the plasma etching process

in a plasma chamber, the reaction space comprising the plasma chamber.

6. (Previously Presented) The method of Claim 5, wherein the exposing the

planarized insulating material to the plasma etching process comprises exposing the planarized

insulating material to a reactive ion etching process.

7. (Previously Presented) The method of Claim 5, wherein the exposing the

planarized insulating material to the plasma etching process in the plasma chamber comprises

using nitrogen gas.

8. (Previously Presented) The method of Claim 2, wherein the terminating

comprises exposing the planarized insulating material to a nitrogen-containing solution after the

etching.

Filing Date: December 2, 2015

9. (Previously Presented) The method of Claim 8, wherein the nitrogen-containing solution comprises an ammonia-based solution.

- 10. (Previously Presented) The method of Claim 2, wherein the first material comprises a second planarized insulating material on a second semiconductor wafer.
- 11. (Previously Presented) The method of Claim 10, wherein the first material comprises silicon oxide.
- 12. (Previously Presented) The method of Claim 2, wherein the bringing into direct contact comprises bringing into direct contact the terminated planarized insulating material and a silicon material having a native oxide, the first material comprising the silicon material.
- 13. (Previously Presented) The method of Claim 2, further comprising, after the terminating, rinsing the insulating material.
- 14. (Previously Presented) The method of Claim 13, wherein the rinsing comprises immersing the insulating material in deionized water.
- 15. (Previously Presented) The method of Claim 2, wherein the bringing into direct contact is performed at room temperature.
- 16. (Previously Presented) The method of Claim 2, wherein the bringing into direct contact is conducted in air.
- 17. (Previously Presented) The method of Claim 2, wherein the forming the planarized insulating material comprises forming silicon oxide on the semiconductor wafer.
- 18. (Previously Presented) The method of Claim 2, further comprising forming a bond between the planarized insulating material and the first material with a strength of at least 500 mJ/m^2 without annealing at more than about 200° C.
- 19. (Previously Presented) The method of Claim 2, further comprising forming a bond between the planarized insulating material and the first material with a strength of at least 2000 mJ/m^2 without annealing at more than about 200° C.
- 20. (Previously Presented) The method of Claim 2, further comprising annealing the planarized insulating material and the first material at a temperature of no more than about 200° C after the bringing into direct contact.
- 21. (Previously Presented) The method of Claim 2, wherein the planarized insulating material has a surface roughness between about 0.5 and 1.5 nm after the etching and before the bringing into direct contact.

Filing Date: December 2, 2015

22. (Previously Presented) A bonded structure comprising:

a first semiconductor material comprising a planarized insulating material having a first bonding surface; and

a second material having a second bonding surface,

wherein the first bonding surface is an etched surface terminated with a nitrogencontaining species, and

wherein the first and second bonding surfaces are in direct contact with each other and bonded together with a chemical bond without any intervening adhesive.

- 23. (Previously Presented) The structure of Claim 22, further comprising at least one integrated circuit in the first semiconductor material.
- 24. (Previously Presented) The structure of Claim 22, wherein the second material comprises silicon.
- 25. (Previously Presented) The structure of Claim 22, wherein the chemical bond has a strength of at least 500 mJ/m^2 .
- 26. (Previously Presented) The structure of Claim 22, wherein the chemical bond has a strength of at least 2000 mJ/m^2 .
- 27. (Previously Presented) The structure of Claim 22, wherein the planarized insulating material comprises silicon oxide.
- 28. (Previously Presented) The structure of Claim 22, wherein the second bonding surface comprises an etched surface.
- 29. (Previously Presented) The structure of Claim 28, wherein the etched surface of the second bonding surface is terminated with a nitrogen-containing species.
- 30. (Previously Presented) The structure of Claim 22, wherein the first bonding surface is terminated with a nitrogen-containing species by at least one of a nitrogen-containing plasma during etching and a nitrogen-containing solution after etching.
- 31. (Previously Presented) The structure of Claim 22, wherein the chemical bond is a covalent bond.
 - 32. (Previously Presented) A bonding method, comprising: forming a planarized insulating material on a semiconductor wafer; etching the planarized insulating material;

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terminating the planarized insulating material with a nitrogen-containing species by at least one of:

a nitrogen-containing plasma during the etching, and

a nitrogen-containing solution after the etching;

bringing a first material into direct contact with the terminated and planarized insulating material; and

forming a chemical bond between the terminated and planarized insulating material and the first material.

33. (Previously Presented) A bonding method, comprising:

forming a planarized insulating material on a processed semiconductor wafer;

exposing the planarized insulating material to a plasma process in a plasma chamber using a nitrogen-containing gas;

terminating the planarized insulating material with a species;

bringing into direct contact the terminated and planarized insulating material and a first material outside of said plasma chamber; and

forming a chemical bond between the terminated and planarized insulating material and the first material.

34. (Previously Presented) The bonding method of Claim 33, wherein the species comprises nitrogen.

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REMARKS

By way of summary, Claims 2-34 were pending as of the outstanding Office Action. Claims 2-34 stand rejected. No claims are amended herein. Applicant respectfully requests reconsideration in view of the remarks set forth herein.

Claims 22-31 stand rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over Claims 1 and 6 of U.S. Patent Publication No. US 2015/0303263, which corresponds to U.S. Patent Application No. 14/754,111 (hereinafter "the '111 Application"). Claims 22-31 also stand rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over Claim 30 of the '111 Application. Claims 22-31 stand rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over Claims 31 and 36 of the '111 Application. Claims 22-31 stand rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over Claim 60 of the '111 Application.

Claims 22-31 stand rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over Claims 77 and 82 of U.S. Patent No. 8,153,505 (hereinafter "the '505 Patent"). Claims 22-31 stand rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over Claim 115 of the '505 Patent. Claims 22-31 also stand rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over Claims 192 and 197 of the '505 Patent. Claims 22-31 stand rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over Claim 230 of the '505 Patent.

Claims 2-21 and 32-34 stand rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over Claims 1 and 9 of U.S. Patent No. 9,082,627 (hereinafter "the '627 Patent"). Claims 2-21 and 32-34 stand rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over Claims 1, 2, and 11 of the '627 Patent. Note that there appears to be an erroneous reference to the '505 patent in this rejection.

Claims 2-21 and 32-34 stand rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over Claims 39, 40, and 48 of the '505 Patent. Claims 2-21 and 32-34 stand rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over Claims 116, 117, and 126 of the '505 Patent. Claims 2-21 and 32-34

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stand rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over Claims 154, 155, and 163 of the '505 Patent.

Claims 2-21 and 32-34 stand rejected on the ground of nonstatutory obviousness-type double patenting as being unpatentable over Claims 105, 106, and 107 of U.S. Patent 6,902,987 (hereinafter "the '987 Patent").

Without acquiescing to the merits of the double patenting rejections, Applicant files herewith by electronic submission a Terminal Disclaimer for each of the '111 Application, the '505 Patent, the '627 Patent, and the '987 Patent, which disclaims, except as provided in the Terminal Disclaimer, the terminal part of the statutory term of any patent granted on the instant application which would extend beyond the expiration date of the full statutory term of each of the aforementioned patents and patent application.

Furthermore, as M.P.E.P. § 804.02(II) instructs, "[t]he filing of a terminal disclaimer to obviate a rejection based on nonstatutory double patenting is not an admission of the propriety of the rejection. *Quad Environmental Technologies Corp. v. Union Sanitary District*, 946 F.2d 870, 20 USPQ2d 1392 (Fed. Cir. 1991). In *Quad Environmental Technologies*, The court indicated that the 'filing of a terminal disclaimer simply serves the statutory function of removing the rejection of double patenting, and raises neither a presumption nor estoppel on the merits of the rejection."

Accordingly, Applicant respectfully submits that the outstanding double patenting rejections are most and that the Application is in condition for allowance.

No Disclaimers or Disavowals

Although the present communication may include alterations to the application or claims, or characterizations of claim scope or referenced art, Applicant is not conceding in this application that previously pending claims are not patentable over the cited references. Rather, any alterations or characterizations are being made to facilitate expeditious prosecution of this application. Applicant reserves the right to pursue at a later date any previously pending or other broader or narrower claims that capture any subject matter supported by the present disclosure, including subject matter found to be specifically disclaimed herein or by any prior prosecution. Accordingly, reviewers of this or any parent, child or related prosecution history shall not

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reasonably infer that Applicant has made any disclaimers or disavowals of any subject matter supported by the present application.

Co-Pending Applications of Assignee

Applicant wishes to draw the Examiner's attention to the following co-pending applications of the present application's assignee.

Docket No.	Serial No.	Title	Filed
TSSRA.012D2	14/754,111	METHOD FOR LOW TEMPERATURE	June 29,
	14//34,111	BONDING AND BONDED STRUCTURE	2015
TSSRA.015C3	14/746,425	THREE DIMENSIONAL DEVICE INTEGRATION METHOD AND INTEGRATED DEVICE	June 22, 2015
TSSRA.016C4	14/474,501	METHOD OF ROOM TEMPERATURE COVALENT BONDING	September 2, 2014

Please charge any additional fees, including any fees for additional extension of time, or credit overpayment to Deposit Account No. 11-1410.

Respectfully submitted,

KNOBBE, MARTENS, OLSON & BEAR, LLP

Dated: February 8, 2016 By: /Paul Stellman/

Paul Stellman Registration No. 68,287 Attorney of Record Customer No. 135,980 (415) 954-4114

22646757 020816

Electronic Acknowledgement Receipt			
EFS ID:	24856867		
Application Number:	14957501		
International Application Number:			
Confirmation Number:	1033		
Title of Invention:	METHOD FOR LOW TEMPERATURE BONDING AND BONDED STRUCTURE		
First Named Inventor/Applicant Name:	Qin-Yi Tong		
Customer Number:	135980		
Filer:	Paul Steven Stellman/Anthony Bonilla		
Filer Authorized By:	Paul Steven Stellman		
Attorney Docket Number:	TSSRA.012C1		
Receipt Date:	08-FEB-2016		
Filing Date:	02-DEC-2015		
Time Stamp:	18:19:22		
Application Type:	Utility under 35 USC 111(a)		

Payment information:

Submitted with Payment no					
File Listing:					
Document Number	Document Description	File Name	File Size(Bytes)/ Message Digest	Multi Part /.zip	Pages (if appl.)
1		2016-02-08_Response_TSSRA-0	135954	yes	8
' 12C1.pdf	3ecfd3a7a3cec193b4632c19aa7adf6fb7a3 973e	· · · · · · · · · · · · · · · · · · ·	J		

	Multipart Description/PDF files in .zip description		
	Document Description	Start	End
	Amendment/Req. Reconsideration-After Non-Final Reject	1	1
	Claims	2	5
	Applicant Arguments/Remarks Made in an Amendment	6	8
•		•	

Warnings:

Information:

Total Files Size (in bytes):	135954

This Acknowledgement Receipt evidences receipt on the noted date by the USPTO of the indicated documents, characterized by the applicant, and including page counts, where applicable. It serves as evidence of receipt similar to a Post Card, as described in MPEP 503.

New Applications Under 35 U.S.C. 111

If a new application is being filed and the application includes the necessary components for a filing date (see 37 CFR 1.53(b)-(d) and MPEP 506), a Filing Receipt (37 CFR 1.54) will be issued in due course and the date shown on this Acknowledgement Receipt will establish the filing date of the application.

National Stage of an International Application under 35 U.S.C. 371

If a timely submission to enter the national stage of an international application is compliant with the conditions of 35 U.S.C. 371 and other applicable requirements a Form PCT/DO/EO/903 indicating acceptance of the application as a national stage submission under 35 U.S.C. 371 will be issued in addition to the Filing Receipt, in due course.

New International Application Filed with the USPTO as a Receiving Office

If a new international application is being filed and the international application includes the necessary components for an international filing date (see PCT Article 11 and MPEP 1810), a Notification of the International Application Number and of the International Filing Date (Form PCT/RO/105) will be issued in due course, subject to prescriptions concerning national security, and the date shown on this Acknowledgement Receipt will establish the international filing date of the application.

	Application No.	14/957,501
INFORMATION DISCLOSURE	Filing Date	December 2, 2015
STATEMENT BY APPLICANT	First Named Inventor	Tong et al.
STATEMENT BY APPLICANT	Art Unit	2895
(Multiple sheets used when necessary)	Examiner	Joannie A. Garcia
SHEET 1 OF 22	Attorney Docket No.	TSSRA.012C1

			U.S. PATENT	DOCUMENTS	
Examiner Initials	Cite No.	Document Number Number - Kind Code (if known) Example: 1,234,567 B1	Publication Date MM-DD-YYYY	Name	Pages, Columns, Lines Where Relevant Passages or Relevant Figures Appear
	1	3,423,823	01-28-1969	Ansley	
	2	3,488,834	01-13-1970	Baird	
	3	3,508,980	04-28-1970	Jackson, Jr. et al.	
	4	3,534,467	10-20-1970	Sachs et al.	
	5	3,579,391	05-18-1971	Buie	
	6	3,587,166	06-28-1971	Alexander et al.	
	7	3,602,981	09-07-1971	Kool	
	8	3,607,466	09-21-1971	Miyazaki	
	9	3,640,807	02-08-1972	Van Dijk	
	10	3,647,581	03-07-1972	Mash	
	11	3,888,708	06-10-1975	Wise et al.	
	12	4,416,054	11-22-1983	Thomas et al.	
	13	4,612,083	09-16-1986	Yasumoto et al.	
	14	4,617,160	10-14-1986	Belanger et al.	
	15	4,649,630	03-13-1987	Boland et al.	
	16	4,700,466	10-20-1987	Nakagawa et al.	
	17	4,754,544	07-05-1988	Hanak	
	18	4,829,018	05-09-1989	Wahlstrom	
	19	4,935,386	06-19-1990	Nakagawa	
	20	4,939,101	07-03-1990	Black et al.	
	21	4,962,879	10-16-1990	Gösele et al.	
	22	4,963,505	10-16-1990	Fujii et al.	
	23	4,970,175	11-13-1990	Haisma et al.	
	24	4,978,421	12-18-1990	Bassous et al.	
	25	4,983,251	01-08-1991	Haisma et al.	
	26	4,992,149	02-12-1991	Nguyen	
	27	5,007,071	04-09-1991	Nakano	
	28	5,013,471	05-07-1991	Ogawa	
	29	5,024,723	06-18-1991	Gösele et al.	

Examiner Signature	Date Considered

^{*}Examiner: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

	Application No.	14/957,501
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STATEMENT BY APPLICANT	First Named Inventor	Tong et al.
STATEMENT BY ALL LICANT	Art Unit	2895
(Multiple sheets used when necessary)	Examiner	Joannie A. Garcia
SHEET 2 OF 22	Attorney Docket No.	TSSRA.012C1

			U.S. PATENT	DOCUMENTS	
Examiner Initials	Cite No.	Document Number Number - Kind Code (if known) Example: 1,234,567 B1	Publication Date MM-DD-YYYY	Name	Pages, Columns, Lines Where Relevant Passages or Relevant Figures Appear
	30	5,034,343	07-23-1991	Rouse et al.	
	31	5,070,026	12-03-1991	Greenwald et al.	
	32	5,071,792	12-10-1991	Van Vonno et al.	
	33	5,081,061	01-14-1992	Rouse et al.	
	34	5,087,585	02-11-1992	Hayashi	
	35	5,089,431	02-18-1992	Slatter et al.	
	36	5,121,706	06-16-1992	Nichols et al.	
	37	5,162,251	11-10-1992	Poole et al.	
	38	5,183,783	02-02-1993	Ohta et al.	
	39	5,196,375	03-23-1993	Hoshi	
	40	5,207,864	05-04-1993	Bhat et al.	
	41	5,234,860	08-10-1993	Gluck	
	42	5,236,118	08-17-1993	Bower et al.	
	43	5,238,875	08-24-1993	Ogino	
	44	5,266,511	11-30-1993	Takao	
	45	5,266,824	11-30-1993	Abe	
	46	5,270,259	12-14-1993	Ito et al.	
	47	5,270,261	12-14-1993	Bertin et al.	
	48	5,272,104	12-21-1993	Schrantz et al.	
	49	5,284,707	02-08-1994	Ogawa et al.	
	50	5,321,301	06-14-1994	Sato et al.	
	51	5,324,566	06-28-1994	Ogawa et al.	
	52	5,324,687	06-28-1994	Wojnarowski	
	53	5,354,695	10-11-1994	Leedy	
	54	5,362,659	11-08-1994	Cartagena	
	55	5,376,579	12-27-1994	Annamalai	
	56	5,380,681	01-10-1995	Hsu	
	57	5,383,993	01-24-1995	Katada et al.	
	58	5,407,856	04-18-1995	Quenzer et al.	

Examiner Signature	Date Considered
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STATEMENT BY APPLICANT	Art Unit	2895
(Multiple sheets used when necessary)	Examiner	Joannie A. Garcia
SHEET 3 OF 22	Attorney Docket No.	TSSRA.012C1

			U.S. PATENT	DOCUMENTS	
Examiner Initials	Cite No.	Document Number Number - Kind Code (if known) Example: 1,234,567 B1	Publication Date MM-DD-YYYY	Name	Pages, Columns, Lines Where Relevant Passages or Relevant Figures Appear
	59	5,413,955	05-09-1995	Lee et al.	
	60	5,420,449	05-30-1995	Oji	
	61	5,421,953	06-06-1995	Nagakubo et al.	
	62	5,422,302	06-06-1995	Yonehara	
	63	5,427,638	06-27-1995	Goetz et al.	
	64	5,432,729	07-11-1995	Carson et al.	
	65	5,437,894	08-01-1995	Ogawa et al.	
	66	5,441,591	08-15-1995	Imthum et al.	
	67	5,441,911	08-15-1995	Malhi	
	68	5,459,104	10-17-1995	Sakai	
	69	5,460,659	10-24-1995	Krut	
	70	5,485,540	01-16-1996	Eda	
	71	5,489,554	02-06-1996	Gates	
	72	5,497,033	03-05-1996	Fillion et al.	
	73	5,503,704	04-02-1996	Bower et al.	
	74	5,514,235	05-07-1996	Mitani et al.	
	75	5,516,727	05-14-1996	Broom	
	76	5,517,754	05-21-1996	Bellstein, Jr., et. al.	
	77	5,523,602	06-04-1996	Horiuchi	
	78	5,534,465	07-09-1996	Frye et al.	
	79	5,543,648	08-06-1996	Miyawaki	
	80	5,546,494	08-13-1996	Eda	
	81	5,547,896	08-20-1996	Linn et al.	
	82	5,548,178	08-20-1996	Eda et al.	
	83	5,561,303	10-01-1996	Schrantz et al.	
	84	5,563,084	10-08-1996	Ramm et al.	
	85	5,567,657	10-22-1996	Wonjnarowski et al.	
	86	5,569,620	10-29-1996	Linn et al.	
	87	5,580,407	12-03-1996	Haisma et al.	

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(Multiple sheets used when necessary)	Examiner	Joannie A. Garcia
SHEET 4 OF 22	Attorney Docket No.	TSSRA.012C1

U.S. PATENT DOCUMENTS					
Examiner Initials	Cite No.	Document Number Number - Kind Code (if known) Example: 1,234,567 B1	Publication Date MM-DD-YYYY	Name	Pages, Columns, Lines Where Relevant Passages or Relevant Figures Appear
	88	5,591,678	01-07-1997	Bendik et al.	
	89	5,627,106	05-06-1997	Hsu	
	90	5,647,932	07-15-1997	Taguchi et al.	
	91	5,650,353	07-22-1997	Yoshizawa et al.	
	92	5,652,436	07-29-1997	Stoner et al.	
	93	5,653,019	08-05-1997	Bernhardt et al.	
	94	5,659,192	08-19-1997	Sarma et al.	
	95	5,661,316	08-26-1997	Kish, Jr., et al.	
	96	5,661,901	09-02-1997	King	
	97	5,666,706	09-16-1997	Tomita et al.	
	98	5,668,057	09-16-1997	Eda et al.	
	99	5,672,240	09-30-1997	Stoner et al.	
	100	5,673,478	10-07-1997	Beene et al.	
	101	5,698,471	12-16-1997	Namba et al.	
	102	5,741,733	04-21-1998	Bertagnolli et al.	
	103	5,747,857	05-05-1998	Eda et al.	
	104	5,753,529	05-19-1998	Chang et al.	
	105	5,755,914	05-26-1998	Yonehara	
	106	5,759,753	06-02-1998	Namba et al.	
	107	5,760,478	06-02-1998	Bazso et al.	
	108	5,763,318	06-09-1998	Bertin et al.	
	109	5,766,984	06-16-1998	Ramm et al.	
	110	5,771,555	06-30-1998	Eda et al.	
	111	5,783,477	07-21-1998	Kish, Jr. et al.	
	112	5,785,874	07-28-1998	Eda et al.	
	113	5,793,115	08-11-1998	Zavracky et al.	
	114	5,804,086	09-08-1998	Bruel	
	115	5,807,783	09-15-1998	Gaul et al.	
	116	5,821,665	10-13-1998	Onishi et al.	

Examiner Signature	Date Considered
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	Application No.	14/957,501
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STATEMENT BY APPLICANT	Art Unit	2895
(Multiple sheets used when necessary)	Examiner	Joannie A. Garcia
SHEET 5 OF 22	Attorney Docket No.	TSSRA.012C1

			U.S. PATENT	DOCUMENTS	
Examiner Initials	Cite No.	Document Number Number - Kind Code (if known) Example: 1,234,567 B1	Publication Date MM-DD-YYYY	Name	Pages, Columns, Lines Where Relevant Passages or Relevant Figures Appear
	117	5,841,197	11-24-1998	Adamic, Jr.	
	118	5,849,627	12-15-1998	Linn et al.	
	119	5,851,894	12-22-1998	Ramm	
	120	5,858,876	01-12-1999	Chew	
	121	5,866,942	02-02-1999	Suzuki et al.	
	122	5,869,354	02-09-1999	Leedy et al.	
	123	5,872,025	02-16-1999	Cronin et al.	
	124	5,876,497	03-02-1999	Atoji	
	125	5,877,034	03-02-1999	Ramm et al.	
	126	5,877,070	03-02-1999	Gösele et al.	
	127	5,877,516	03-02-1999	Mermagen et al.	
	128	5,880,010	03-09-1999	Davidson	
	129	5,889,302	03-30-1999	Liu	
	130	5,902,118	05-11-1999	Hubner	
	131	5,903,018	05-11-1999	Shimawaki	
	132	5,904,860	05-18-1999	Nagakubo et al.	
	133	5,910,669	06-08-1999	Namba et al.	
	134	5,915,167	06-22-1999	Leedy	
	135	5,915,193	06-22-1999	Tong et al	
	136	5,920,142	07-06-1999	Onishi et al.	
	137	5,936,280	08-10-1999	Liu	
	138	5,966,622	10-12-1999	Levine et al.	
	139	5,980,770	11-09-1999	Ramachandran et al.	
	140	5,982,010	11-09-1999	Namba et al.	
	141	5,990,472	11-23-1999	Rinne	
	142	5,990,562	11-23-1999	Vallett	
	143	5,991,989	11-30-1999	Onishi et al.	
	144	6,004,865	12-21-1999	Horiuchi	
	145	6,004,866	12-21-1999	Nakano et al.	

Examiner Signature	Date Considered
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	Application No.	14/957,501
INFORMATION DISCLOSURE	Filing Date	December 2, 2015
STATEMENT BY APPLICANT	First Named Inventor	Tong et al.
STATEMENT BY AFFLICANT	Art Unit	2895
(Multiple sheets used when necessary)	Examiner	Joannie A. Garcia
SHEET 6 OF 22	Attorney Docket No.	TSSRA.012C1

			U.S. PATENT	DOCUMENTS	
Examiner Initials	Cite No.	Document Number Number - Kind Code (if known) Example: 1,234,567 B1	Publication Date MM-DD-YYYY	Name	Pages, Columns, Lines Where Relevant Passages or Relevant Figures Appear
	146	6,018,211	01-25-2000	Kanaboshi et al.	
	147	6,048,752	04-11-2000	Lindemann	
	148	6,054,371	09-29-1998	Tsuchiaki et al.	
	149	6,071,761	06-06-2000	Jacobs	
	150	6,087,719	04-23-1998	Tsunashima	
	151	6,087,760	07-11-2000	Yamaguchi et al.	
	152	6,103,009	08-15-2000	Atoji	
	153	6,120,917	09-19-2000	Eda	
	154	6,133,640	10-17-2000	Leedy	
	155	6,136,691	05-26-1998	Chen	
	156	6,143,628	11-07-2000	Sato et al.	
	157	6,146,992	11-14-2000	Lauterbach et al.	
	158	6,153,445	11-28-2000	Yamazaki et al.	
	159	6,153,495	11-28-2000	Kub et al.	
	160	6,154,940	12-05-2000	Onishi et al.	
	161	6,156,624	12-05-2000	Yamagata et al.	
	162	6,165,817	12-26-2000	Akram et al.	
	163	6,180,496	01-30-2001	Farrens et al.	
	164	6,190,778	02-20-2001	Batz-Sohn et al.	
	165	6,194,323	02-27-2001	Downey et al.	
	166	6,197,663	03-06-2001	Chandross et al.	
	167	6,198,159	03-06-2001	Hosoma et al.	
	168	6,236,141	05-22-2001	Sato et al.	
	169	6,242,324	06-05-2001	Kub	
	170	6,246,068	06-12-2001	Sato et al.	
	171	6,255,731	07-03-2001	Ohmi et al.	
	172	6,255,899	07-03-2001	Bertin et al.	
	173	6,270,202	08-07-2001	Namba et al.	
	174	6,274,892	08-14-2001	Kub et al.	

Examiner Signature	Date Considered

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STATEMENT BY ALL LICANT	Art Unit	2895
(Multiple sheets used when necessary)	Examiner	Joannie A. Garcia
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STATEMENT BY AFFEIGANT	Art Unit	2895
(Multiple sheets used when necessary)	Examiner	Joannie A. Garcia
SHEET 15 OF 22	Attorney Docket No.	TSSRA.012C1

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Examiner Signature	Date Considered

^{*}Examiner: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

	Application No.	14/957,501
INFORMATION DISCLOSURE	Filing Date	December 2, 2015
STATEMENT BY APPLICANT	First Named Inventor	Tong et al.
STATEMENT BY AFFLICANT	Art Unit	2895
(Multiple sheets used when necessary)	Examiner	Joannie A. Garcia
SHEET 21 OF 22	Attorney Docket No.	TSSRA.012C1

		NON PATENT LITERATURE DOCUMENTS	
Examiner Initials	Cite No.	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T ¹
	440	U. Gösele et al., History and Future of Semiconductor Wafer Bonding, 47-48 Solid State Phenomena 33 (1996).	
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Examiner Signature	Date Considered
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	Application No.	14/957,501
INFORMATION DISCLOSURE	Filing Date	December 2, 2015
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STATEMENT BY ALL LICANT	Art Unit	2895
(Multiple sheets used when necessary)	Examiner	Joannie A. Garcia
SHEET 22 OF 22	Attorney Docket No.	TSSRA.012C1

		NON PATENT LITERATURE DOCUMENTS	
Examiner Initials Cite No. Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.			
	459	Zucker et al., "Application of Oxygen Plasma Processing to Silicon Direct Bonding", Sensors and Actuators A, 36 (1993), pp. 227-231.	
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Electronic Acknowledgement Receipt		
EFS ID:	24920204	
Application Number:	14957501	
International Application Number:		
Confirmation Number:	1033	
Title of Invention:	METHOD FOR LOW TEMPERATURE BONDING AND BONDED STRUCTURE	
First Named Inventor/Applicant Name:	Qin-Yi Tong	
Customer Number:	135980	
Filer:	Paul Steven Stellman/Jonathan Pan	
Filer Authorized By:	Paul Steven Stellman	
Attorney Docket Number:	TSSRA.012C1	
Receipt Date:	16-FEB-2016	
Filing Date:	02-DEC-2015	
Time Stamp:	13:28:02	
Application Type:	Utility under 35 USC 111(a)	

Payment information:

Submitted with Payment	no
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File Listing:

Number	Document Description	File Name	File Size(Bytes)/ Message Digest	Multi Part /.zip	Pages (if appl.)
1	Non Patent Literature	Plossl-et-al_Wafer-direct- bonding_tailoring-adhesion- between-brittle-materials.pdf	3587507 62/5e1cff280c536f54c5e49fcfcfe46fea871b 4	no	88

Informations

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New Applications Under 35 U.S.C. 111

If a new application is being filed and the application includes the necessary components for a filing date (see 37 CFR 1.53(b)-(d) and MPEP 506), a Filing Receipt (37 CFR 1.54) will be issued in due course and the date shown on this Acknowledgement Receipt will establish the filing date of the application.

National Stage of an International Application under 35 U.S.C. 371

If a timely submission to enter the national stage of an international application is compliant with the conditions of 35 U.S.C. 371 and other applicable requirements a Form PCT/DO/EO/903 indicating acceptance of the application as a national stage submission under 35 U.S.C. 371 will be issued in addition to the Filing Receipt, in due course.

New International Application Filed with the USPTO as a Receiving Office

If a new international application is being filed and the international application includes the necessary components for an international filing date (see PCT Article 11 and MPEP 1810), a Notification of the International Application Number and of the International Filing Date (Form PCT/RO/105) will be issued in due course, subject to prescriptions concerning national security, and the date shown on this Acknowledgement Receipt will establish the international filing date of the application.

Docket No.: TSSRA.012C1 Customer No. 135,980

INFORMATION DISCLOSURE STATEMENT

Inventor : Tong et al.

App. No. : 14/957,501

Filed : December 2, 2015

For : METHOD FOR LOW

TEMPERATURE BONDING AND

BONDED STRUCTURE

Examiner : Joannie A. Garcia

Art Unit : 2895

Conf. No. : 1033

Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450

References and Listing

Pursuant to 37 CFR 1.56, an Information Disclosure Statement listing references is provided herewith. Listed references are of record in one or more of U.S. patent application No. 14/754,111, filed June 29, 2015, U.S. patent application No. 14/197,070, filed March 4, 2014, U.S. patent application No. 13/341,273, filed December 30, 2011, U.S. patent application No. 12/954,740, filed November 26, 2010, U.S. patent application No. 12/720,368, filed March 9, 2010, U.S. patent application No. 11/980,664, filed October 31, 2007, U.S. patent application No. 10/913,441, filed August 9, 2004, and U.S. patent application No. 09/505,283, filed February 16, 2000, which are parent applications of this continuation application, and are relied upon for an earlier filing date under 35 USC 120. Copies of the references are not submitted pursuant to 37 CFR 1.98(d).

No Disclaimers

To the extent that anything in the Information Disclosure Statement or the listed references could be construed as a disclaimer of any subject matter supported by the present application, Applicant hereby rescinds and retracts such disclaimer.

Application No.: 14/957,501

Filing Date: December 2, 2015

Timing of Disclosure

This Information Disclosure Statement is being filed within three months of the filing date, and no fee is believed to be required.

The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment, to Account No. 11-1410.

Respectfully submitted,

KNOBBE, MARTENS, OLSON & BEAR, LLP

Dated: February 16, 2016 By: /Paul Stellman/

Paul Stellman Registration No. 68,287 Attorney of Record Customer No. 135,980 (415) 954-4114

22625377 021616 Docket No.: TSSRA.012C1 February 29, 2016
Page 1 of 1

Please Direct All Correspondence to Customer Number 135,980

RESPONSE TO INFORMATIONAL NOTICE

Inventor : Tong et al.

App. No. : 14/957,501

Filed : December 2, 2015

For : METHOD FOR LOW

TEMPERATURE BONDING AND

BONDED STRUCTURE

Art Unit : 2895

Conf No. : 1033

Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450

Dear Sir:

The above-captioned application was filed without a Declaration and/or Substitute Statement. Enclosed in compliance with 37 CFR 1.53(f) are the following.

(X) Substitute Statement in Lieu of Declaration document(s) for:

Qin-Yi Tong

The Commissioner is hereby authorized to charge any additional fees which may be required, now or in the future, or credit any overpayment, to Account No. 11-1410.

/Paul Stellman/

Paul Stellman Registration No. 68,287 Attorney of Record Customer No. 135,980 (415) 954-4114

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SUBSTITUTE STATEMENT IN LIEU OF AN OATH OR DECLARATION FOR UTILITY OR DESIGN PATENT APPLICATION (35 U.S.C. 115(d) AND 37 CFR 1.64)

Title of Invention	METHOD FOR LOW TEMP	ERATURE BONDIN	G AND BONDED ST	RUCTURE
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I believe the above-named inventor or joint inventor to be the original inventor or an original joint inventor of a claimed inventon in the application. The above-identified application was made or authorized to be made by me. I hereby acknowledge that any willful false statement made in this statement is punishable under 18 U.S.C. 1001 by fine or imprisonment of not more than five (5) years, or both.				
Relationship to the inventor to whom this substitute statement applies: Legal Representative (for deceased or legally incapacitated inventor only), Assignee, Person to whom the inventor is under an obligation to assign, Person who otherwise shows a sufficient proprietary interest in the matter (petition under 37 CFR 1.48 is required), or Joint Inventor.				

[Page 1 of 2]

This collection of information is required by 35 U.S.C. 115 and 37 CFR 1.93. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.11 and 1.14. This collection is estimated to take 1 minute to complete, including gethering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Accomments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief information Officer, U.S. Patent and Trademark Office, U.S. Department of Commence, P.O. Box 1450, Alexandria, VA 22313-1450, DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS, SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.

SUBSTITUTE STATEMENT

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Circumstances permitting execution	of this substitute statemen	it:		
Inventor is deceased,				
Inventor is under legal ince	pacity,			
inventor cannot be found or	r reached after diligent effo	m, or		
inventor has refused to exe	oute the eath or declaratio	n under 37 CFR 1,60	3.	
If there are joint inventors, please of	eck the appropriate box br	eiow:		
An application data sheet u or is currently submitted.	nder 37 CFR 1.76 (PTO/A)	IA/14 or equivalent) i	naming the entire	inventive entity has been
OR 				
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PERSON EXECUTING THIS SUBSTIT	UTE STATEMENT:			
Name: Paul Davis				1/6/2016 ale (Optional):
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APPLICANT NAME AND TITLE OF PI	ERSON EXECUTING THIS	3 SUBSTITUTE STA	TEMENT:	
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Electronic Acknowledgement Receipt			
EFS ID:	25052671		
Application Number:	14957501		
International Application Number:			
Confirmation Number:	1033		
Title of Invention:	METHOD FOR LOW TEMPERATURE BONDING AND BONDED STRUCTURE		
First Named Inventor/Applicant Name:	Qin-Yi Tong		
Customer Number:	135980		
Filer:	Paul Steven Stellman/Kealani Aguon		
Filer Authorized By:	Paul Steven Stellman		
Attorney Docket Number:	TSSRA.012C1		
Receipt Date:	29-FEB-2016		
Filing Date:	02-DEC-2015		
Time Stamp:	17:15:33		
Application Type:	Utility under 35 USC 111(a)		

Payment information:

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File Listing:					
Document Number	Document Description	File Name	File Size(Bytes)/ Message Digest	Multi Part /.zip	Pages (if appl.)
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New Applications Under 35 U.S.C. 111

If a new application is being filed and the application includes the necessary components for a filing date (see 37 CFR 1.53(b)-(d) and MPEP 506), a Filing Receipt (37 CFR 1.54) will be issued in due course and the date shown on this Acknowledgement Receipt will establish the filing date of the application.

National Stage of an International Application under 35 U.S.C. 371

If a timely submission to enter the national stage of an international application is compliant with the conditions of 35 U.S.C. 371 and other applicable requirements a Form PCT/DO/EO/903 indicating acceptance of the application as a national stage submission under 35 U.S.C. 371 will be issued in addition to the Filing Receipt, in due course.

New International Application Filed with the USPTO as a Receiving Office

If a new international application is being filed and the international application includes the necessary components for an international filing date (see PCT Article 11 and MPEP 1810), a Notification of the International Application Number and of the International Filing Date (Form PCT/RO/105) will be issued in due course, subject to prescriptions concerning national security, and the date shown on this Acknowledgement Receipt will establish the international filing date of the application.

	Application No.	14/957,501
INFORMATION DISCLOSURE	Filing Date	December 2, 2015
STATEMENT BY APPLICANT	First Named Inventor	Tong et al.
STATEMENT BY AFFEIGANT	Art Unit	2895
(Multiple sheets used when necessary)	Examiner	Joannie A. Garcia
SHEET 1 OF 1	Attorney Docket No.	TSSRA.012C1

	U.S. PATENT DOCUMENTS				
Examiner Initials	Cite No.	Document Number Number - Kind Code (if known) Example: 1,234,567 B1	Publication Date MM-DD-YYYY	Name	Pages, Columns, Lines Where Relevant Passages or Relevant Figures Appear

FOREIGN PATENT DOCUMENTS						
Examiner Initials	Cite No.	Foreign Patent Document Country Code-Number-Kind Code Example: JP 1234567 A1	Publication Date MM-DD-YYYY	Name	Pages, Columns, Lines Where Relevant Passages or Relevant Figures Appear	T ¹

	NON PATENT LITERATURE DOCUMENTS					
Examiner Initials	Cite No.	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T ¹			
	1	AMIRFEIZ et al., "Formation of silicon structures by plasma activated wafer bonding," Vol. 99-2, Abstract No. 963, 1999 Joint Int'l Meeting of the Electrochecm. Soc'y, December 16, 1999.				
	2	AMIRFEIZ et al., "Formation of silicon structures by plasma activated wafer bonding," Electrochemical Society Proceedings: Semiconductor Wafer Bonding: Science, Technology, and Applications V, Volume 99-35, (1999), pp. 29-39.				
	3	Ziptronix, Inc. v. Omnvision Technologies, Inc., Defendants' Third Supplemental Consolidated Invalidity Contentions Under Patent Local Rule 3-3, Case No. 4:10-cv-05525-SBA, dated June 14, 2013, pp. 1-5379, (submitted in multiple parts).				

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Examiner Signature	Date Considered

^{*}Examiner: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

Electronic Patent Application Fee Transmittal						
Application Number:	14957501					
Filing Date:	02-Dec-2015					
Title of Invention:	METHOD FOR LOW TEMPERATURE BONDING AND BONDED STRUCTURE				DED STRUCTURE	
First Named Inventor/Applicant Name:	Qir	n-Yi Tong				
Filer:	Pau	ul Steven Stellman/.	Jonathan Pan			
Attorney Docket Number:	TSS	SRA.012C1				
Filed as Large Entity						
Filing Fees for Utility under 35 USC 111(a)						
Description		Fee Code	Quantity	Amount	Sub-Total in USD(\$)	
Basic Filing:						
Pages:						
Claims:						
Miscellaneous-Filing:						
Petition:						
Patent-Appeals-and-Interference:						
Post-Allowance-and-Post-Issuance:						
Extension-of-Time:						

Description	Fee Code	Quantity	Amount	Sub-Total in USD(\$)
Miscellaneous:				
Submission- Information Disclosure Stmt	1806	1	180	180
	Tot	al in USD	(\$)	180

Electronic Acknowledgement Receipt				
EFS ID:	25178415			
Application Number:	14957501			
International Application Number:				
Confirmation Number:	1033			
Title of Invention:	METHOD FOR LOW TEMPERATURE BONDING AND BONDED STRUCTURE			
First Named Inventor/Applicant Name:	Qin-Yi Tong			
Customer Number:	135980			
Filer:	Paul Steven Stellman/Jonathan Pan			
Filer Authorized By:	Paul Steven Stellman			
Attorney Docket Number:	TSSRA.012C1			
Receipt Date:	11-MAR-2016			
Filing Date:	02-DEC-2015			
Time Stamp:	19:02:11			
Application Type:	Utility under 35 USC 111(a)			

Payment information:

Submitted with Payment	yes
Payment Type	Credit Card
Payment was successfully received in RAM	\$180
RAM confirmation Number	5259
Deposit Account	111410
Authorized User	KNOBBE MARTENS OLSON AND BEAR

The Director of the USPTO is hereby authorized to charge indicated fees and credit any overpayment as follows:

Charge any Additional Fees required under 37 CFR 1.16 (National application filing, search, and examination fees)

File Listing:									
Document Number	Document Description	File Name	File Size(Bytes)/ Message Digest	Multi Part /.zip	Pages (if appl.)				
1		2016-03-11_IDS_TSSRA-012C1.	144402	yes	3				
·		pdf	9f0024d0219d102a969cf3fd628ac1f7d237 c31d	,	3				
	Multipart Description/PDF files in .zip description								
	Document D	Document Description			nd				
	Transmitta	Transmittal Letter			2				
	Information Disclosure Stat	ement (IDS) Form (SB08)	3	3					
Warnings:									
Information:									
2	Fee Worksheet (SB06) fee-info.pdf		30909	no	2				
		·	ec693c5756dcfa3947854bfee15334358949 561d						
Warnings:									
Information:									
		Total Files Size (in bytes)	17	75311					

This Acknowledgement Receipt evidences receipt on the noted date by the USPTO of the indicated documents, characterized by the applicant, and including page counts, where applicable. It serves as evidence of receipt similar to a Post Card, as described in MPEP 503.

New Applications Under 35 U.S.C. 111

If a new application is being filed and the application includes the necessary components for a filing date (see 37 CFR 1.53(b)-(d) and MPEP 506), a Filing Receipt (37 CFR 1.54) will be issued in due course and the date shown on this Acknowledgement Receipt will establish the filing date of the application.

National Stage of an International Application under 35 U.S.C. 371

If a timely submission to enter the national stage of an international application is compliant with the conditions of 35 U.S.C. 371 and other applicable requirements a Form PCT/DO/EO/903 indicating acceptance of the application as a national stage submission under 35 U.S.C. 371 will be issued in addition to the Filing Receipt, in due course.

New International Application Filed with the USPTO as a Receiving Office

If a new international application is being filed and the international application includes the necessary components for an international filing date (see PCT Article 11 and MPEP 1810), a Notification of the International Application Number and of the International Filing Date (Form PCT/RO/105) will be issued in due course, subject to prescriptions concerning national security, and the date shown on this Acknowledgement Receipt will establish the international filing date of the application.

Docket No.: TSSRA.012C1 Customer No. 135,980

INFORMATION DISCLOSURE STATEMENT

First Inventor: Tong et al.

App. No. : 14/957,501

Filed : December 2, 2015

For : METHOD FOR LOW TEMPERATURE

BONDING AND BONDED STRUCTURE

Examiner : Joannie A. Garcia

Art Unit : 2895

Conf. No. : 1033

Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450

References and Listing

Pursuant to 37 CFR 1.56, an Information Disclosure Statement listing references is provided herewith. Listed references are of record in U.S. patent application No. 14/754,111, filed June 29, 2015, which is the parent of this continuation application, and is relied upon for an earlier filing date under 35 USC 120. Copies of the references are not submitted pursuant to 37 CFR 1.98(d).

No Disclaimers

To the extent that anything in the Information Disclosure Statement or the listed references could be construed as a disclaimer of any subject matter supported by the present application, Applicant hereby rescinds and retracts such disclaimer.

Timing of Disclosure

This Information Disclosure Statement is being filed after receipt of a First Office Action, but before the mailing date of a Final Action and before the mailing date of a Notice of Allowance. This Statement is accompanied by the fees set forth in 37 CFR 1.17(p). The Commissioner is hereby authorized to charge any additional fees which may be required or to credit any overpayment to Account No. 11-1410.

Application No.: 14/957,501

Filing Date: December 2, 2015

Respectfully submitted,

KNOBBE, MARTENS, OLSON & BEAR, LLP

Dated: March 11, 2016 By: /Paul Stellman/

Paul Stellman Registration No. 68,287 Attorney of Record Customer No. 135,980 (415) 954-4114

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United States Patent and Trademark Office

INITED STATES DEPARTMENT OF COMMERCE United States Patent and Trademark Office Address: COMMISSIONER FOR PATENTS P.O. Sox 1450 Alexandria, Virginia 22313-1450 www.uspto.gov

APPLICATION NUMBER

FILING OR 371(C) DATE 12/02/2015

FIRST NAMED APPLICANT Qin-Yi Tong

ATTY. DOCKET NO./TITLE TSSRA.012C1

CONFIRMATION NO. 1033

PUBLICATION NOTICE

135980 Knobbe, Martens, Olson & Bear, LLP 2040 main Street Fourteenth Floor Irvine, CA 92614

14/957,501

Title:METHOD FOR LOW TEMPERATURE BONDING AND BONDED STRUCTURE

Publication No.US-2016-0086913-A1 Publication Date: 03/24/2016

NOTICE OF PUBLICATION OF APPLICATION

The above-identified application will be electronically published as a patent application publication pursuant to 37 CFR 1.211, et seg. The patent application publication number and publication date are set forth above.

The publication may be accessed through the USPTO's publically available Searchable Databases via the Internet at www.uspto.gov. The direct link to access the publication is currently http://www.uspto.gov/patft/.

The publication process established by the Office does not provide for mailing a copy of the publication to applicant. A copy of the publication may be obtained from the Office upon payment of the appropriate fee set forth in 37 CFR 1.19(a)(1). Orders for copies of patent application publications are handled by the USPTO's Office of Public Records. The Office of Public Records can be reached by telephone at (703) 308-9726 or (800) 972-6382. by facsimile at (703) 305-8759, by mail addressed to the United States Patent and Trademark Office, Office of Public Records, Alexandria, VA 22313-1450 or via the Internet.

In addition, information on the status of the application, including the mailing date of Office actions and the dates of receipt of correspondence filed in the Office, may also be accessed via the Internet through the Patent Electronic Business Center at www.uspto.gov using the public side of the Patent Application Information and Retrieval (PAIR) system. The direct link to access this status information is currently http://pair.uspto.gov/. Prior to publication, such status information is confidential and may only be obtained by applicant using the private side of PAIR.

Further assistance in electronically accessing the publication, or about PAIR, is available by calling the Patent Electronic Business Center at 1-866-217-9197.

Office of Data Managment, Application Assistance Unit (571) 272-4000, or (571) 272-4200, or 1-888-786-0101

page 1 of 1

05/24/2016

UNITED STATES DEPARTMENT OF COMMERCE United States Patent and Trademark Office Address: COMMISSIONER FOR PATENTS P.O. Box 1450 Alexandria, Virginia 22313-1450

NOTICE OF ALLOWANCE AND FEE(S) DUE

Knobbe, Martens, Olson & Bear, LLP 2040 main Street Fourteenth Floor Irvine, CA 92614

EXAMINER GARCIA, JOANNIE A

ART UNIT PAPER NUMBER

2895

DATE MAILED: 05/24/2016

APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
14/957,501	12/02/2015	Qin-Yi Tong	TSSRA.012C1	1033

TITLE OF INVENTION: METHOD FOR LOW TEMPERATURE BONDING AND BONDED STRUCTURE

APPLN. TYPE	ENTITY STATUS	ISSUE FEE DUE	PUBLICATION FEE DUE	PREV. PAID ISSUE FEE	TOTAL FEE(S) DUE	DATE DUE
nonprovisional	UNDISCOUNTED	\$960	\$0	\$0	\$960	08/24/2016

THE APPLICATION IDENTIFIED ABOVE HAS BEEN EXAMINED AND IS ALLOWED FOR ISSUANCE AS A PATENT. PROSECUTION ON THE MERITS IS CLOSED. THIS NOTICE OF ALLOWANCE IS NOT A GRANT OF PATENT RIGHTS. THIS APPLICATION IS SUBJECT TO WITHDRAWAL FROM ISSUE AT THE INITIATIVE OF THE OFFICE OR UPON PETITION BY THE APPLICANT. SEE 37 CFR 1.313 AND MPEP 1308.

THE ISSUE FEE AND PUBLICATION FEE (IF REQUIRED) MUST BE PAID WITHIN <u>THREE MONTHS</u> FROM THE MAILING DATE OF THIS NOTICE OR THIS APPLICATION SHALL BE REGARDED AS ABANDONED. <u>THIS STATUTORY PERIOD CANNOT BE EXTENDED.</u> SEE 35 U.S.C. 151. THE ISSUE FEE DUE INDICATED ABOVE DOES NOT REFLECT A CREDIT FOR ANY PREVIOUSLY PAID ISSUE FEE IN THIS APPLICATION. IF AN ISSUE FEE HAS PREVIOUSLY BEEN PAID IN THIS APPLICATION (AS SHOWN ABOVE), THE RETURN OF PART B OF THIS FORM WILL BE CONSIDERED A REQUEST TO REAPPLY THE PREVIOUSLY PAID ISSUE FEE TOWARD THE ISSUE FEE NOW DUE.

HOW TO REPLY TO THIS NOTICE:

I. Review the ENTITY STATUS shown above. If the ENTITY STATUS is shown as SMALL or MICRO, verify whether entitlement to that entity status still applies.

If the ENTITY STATUS is the same as shown above, pay the TOTAL FEE(S) DUE shown above.

If the ENTITY STATUS is changed from that shown above, on PART B - FEE(S) TRANSMITTAL, complete section number 5 titled "Change in Entity Status (from status indicated above)".

For purposes of this notice, small entity fees are 1/2 the amount of undiscounted fees, and micro entity fees are 1/2 the amount of small entity fees

II. PART B - FEE(S) TRANSMITTAL, or its equivalent, must be completed and returned to the United States Patent and Trademark Office (USPTO) with your ISSUE FEE and PUBLICATION FEE (if required). If you are charging the fee(s) to your deposit account, section "4b" of Part B - Fee(s) Transmittal should be completed and an extra copy of the form should be submitted. If an equivalent of Part B is filed, a request to reapply a previously paid issue fee must be clearly made, and delays in processing may occur due to the difficulty in recognizing the paper as an equivalent of Part B.

III. All communications regarding this application must give the application number. Please direct all communications prior to issuance to Mail Stop ISSUE FEE unless advised to the contrary.

IMPORTANT REMINDER: Utility patents issuing on applications filed on or after Dec. 12, 1980 may require payment of maintenance fees. It is patentee's responsibility to ensure timely payment of maintenance fees when due.

PART B - FEE(S) TRANSMITTAL

Complete and send this form, together with applicable fee(s), to: Mail Mail Stop ISSUE FEE

Mail Stop ISSUE FEE Commissioner for Patents P.O. Box 1450 Alexandria, Virginia 22313-1450

Alexandria, Virginia or Fax (571)-273-2885

INSTRUCTIONS: This form should be used for transmitting the ISSUE FEE and PUBLICATION FEE (if required). Blocks 1 through 5 should be completed where appropriate. All further correspondence including the Patent, advance orders and notification of maintenance fees will be mailed to the current correspondence address as indicated unless corrected below or directed otherwise in Block 1, by (a) specifying a new correspondence address; and/or (b) indicating a separate "FEE ADDRESS" for maintenance fee notifications.

maintenance fee notifica		ici wise in Bio	CK 1, by (a) specifying a new c		,		., .	Tate TEE ADDRESS TO
CURRENT CORRESPOND	DENCE ADDRESS (Note: Use B	ock 1 for any chang	ge of address)		Note Fee(pape have	e: A certificate of s) Transmittal. Thi rs. Each additiona its own certificate	mailing s certif l paper of mai	g can only be used for icate cannot be used for such as an assignmen ling or transmission.	r domestic mailings of the or any other accompanying nt or formal drawing, must
135980 7590 05/24/2016 Knobbe, Martens, Olson & Bear, LLP 2040 main Street					I her State addr trans	Cer reby certify that th es Postal Service w essed to the Mail mitted to the USP	tificate is Fee(s vith suf Stop TO (57	of Mailing or Transı 3) Transmittal is being ficient postage for firs ISSUE FEE address 1) 273-2885, on the da	mission g deposited with the United t class mail in an envelope above, or being facsimile te indicated below.
Fourteenth Floo									(Depositor's name)
Irvine, CA 9261	4								(Signature)
									(Date)
APPLICATION NO.	FILING DATE			FIRST NAMED INVEN	NTOR		ATTO:	RNEY DOCKET NO.	CONFIRMATION NO.
14/957,501	12/02/2015	<u> </u>		Qin-Yi Tong				TSSRA.012C1	1033
TITLE OF INVENTION	N: METHOD FOR LOW	TEMPERATU	JRE BONI		STR	UCTURE			
APPLN. TYPE	ENTITY STATUS	ISSUE FEI	E DUE	PUBLICATION FEE I	DUE	PREV. PAID ISSUI	E FEE	TOTAL FEE(S) DUE	DATE DUE
nonprovisional	UNDISCOUNTED	\$96	0	\$0		\$0		\$960	08/24/2016
EXAN	MINER	ART U	NIT	CLASS-SUBCLAS	s				
	OANNIE A	289		257-684000					
<u> </u>	ence address or indication			2. For printing on	the n	atent front page lis	ıf.		
CFR 1.363).				(1) The names of	up to	3 registered paten		eys 1	
Address form PTO/S	oondence address (or Cha B/122) attached.	inge of Corresp	ondence	or agents OR, alte		ely, e firm (having as a	memb	era 2	
"Fee Address" inc PTO/SB/47; Rev 03- Number is required	lication (or "Fee Address 02 or more recent) attach •	" Indication fo ed. Use of a C	rm ustomer	registered attorne 2 registered paten listed, no name wi	y or a t attor	gent) and the nameneys or agents. If	es of up no nam	o to e is 3	
	AND RESIDENCE DATA			•		*			
PLEASE NOTE: Un recordation as set for	less an assignee is ident th in 37 CFR 3.11. Com	ified below, no	o assignee form is NO	data will appear on t T a substitute for filin	the pa	itent. If an assign	ee is id	entified below, the do	ocument has been filed for
(A) NAME OF ASSI				(B) RESIDENCE: (
					_	_			_
Please check the appropriate of the control of the	riate assignee category or	categories (wi	ill not be p	rinted on the patent):	Ц	Individual 🖵 Co	orporati	on or other private gro	oup entity Government
4a. The following fee(s)	are submitted:		4			se first reapply ar	ıy prev	iously paid issue fee s	shown above)
Issue Fee	No small entity discount j	itt . d\		A check is enclo		4 Ec PTO 2020	ia atta	de a d	
	of Copies			☐ Payment by credit card. Form PTO-2038 is attached. ☐ The director is hereby authorized to charge the required fee(s), any deficiency, or credits any					iciency, or credits any
				overpayment, to	Depos	sit Account Numbe	ér	(enclose ar	n extra copy of this form).
5 Change in Entity Sta	itus (from status indicate	d above)							
_ ~ .	ng micro entity status. Se)	NOTE: Absent a val	lid cer	tification of Micro	Entity	Status (see forms PTC	D/SB/15A and 15B), issue
☐ Applicant asserting small entity status. See 37 CFR 1.27				fee payment in the micro entity amount will not be accepted at the risk of application abandonment. NOTE: If the application was previously under micro entity status, checking this box will be taken to be a notification of loss of entitlement to micro entity status.					
Applicant changing to regular undiscounted fee status.				NOTE: Checking this box will be taken to be a notification of loss of entitlement to small or micro entity status, as applicable.					
NOTE: This form must	be signed in accordance v	vith 37 CFR 1.	31 and 1.3	3. See 37 CFR 1.4 for	signa	ture requirements	and cer	tifications.	
Authorized Signature						Date			
Typed or printed nam	ne					Registration N	Го		

Page 274 of 331

Page 2 of 3

SAMSUNG EXHIBIT 1002

PTOL-85 Part B (10-13) Approved for use through 10/31/2013.



UNITED STATES PATENT AND TRADEMARK OFFICE

UNITED STATES DEPARTMENT OF COMMERCE United States Patent and Trademark Office Address: COMMISSIONER FOR PATENTS

P.O. Box 1450 Alexandria, Virginia 22313-1450 www.uspto.gov

APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
14/957,501	12/02/2015	12/02/2015 Qin-Yi Tong		1033
135980 75	590 05/24/2016		EXAM	INER
Knobbe, Martens	, Olson		GARCIA, J	DANNIE A
& Bear, LLP				
2040 main Street			ART UNIT	PAPER NUMBER
Fourteenth Floor			2895	
Irvine, CA 92614			DATE MAILED: 05/24/201	6

Determination of Patent Term Adjustment under 35 U.S.C. 154 (b)

(Applications filed on or after May 29, 2000)

The Office has discontinued providing a Patent Term Adjustment (PTA) calculation with the Notice of Allowance.

Section 1(h)(2) of the AIA Technical Corrections Act amended 35 U.S.C. 154(b)(3)(B)(i) to eliminate the requirement that the Office provide a patent term adjustment determination with the notice of allowance. See Revisions to Patent Term Adjustment, 78 Fed. Reg. 19416, 19417 (Apr. 1, 2013). Therefore, the Office is no longer providing an initial patent term adjustment determination with the notice of allowance. The Office will continue to provide a patent term adjustment determination with the Issue Notification Letter that is mailed to applicant approximately three weeks prior to the issue date of the patent, and will include the patent term adjustment on the patent. Any request for reconsideration of the patent term adjustment determination (or reinstatement of patent term adjustment) should follow the process outlined in 37 CFR 1.705.

Any questions regarding the Patent Term Extension or Adjustment determination should be directed to the Office of Patent Legal Administration at (571)-272-7702. Questions relating to issue and publication fee payments should be directed to the Customer Service Center of the Office of Patent Publication at 1-(888)-786-0101 or (571)-272-4200.

OMB Clearance and PRA Burden Statement for PTOL-85 Part B

The Paperwork Reduction Act (PRA) of 1995 requires Federal agencies to obtain Office of Management and Budget approval before requesting most types of information from the public. When OMB approves an agency request to collect information from the public, OMB (i) provides a valid OMB Control Number and expiration date for the agency to display on the instrument that will be used to collect the information and (ii) requires the agency to inform the public about the OMB Control Number's legal significance in accordance with 5 CFR 1320.5(b).

The information collected by PTOL-85 Part B is required by 37 CFR 1.311. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to take 12 minutes to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, Virginia 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, Virginia 22313-1450. Under the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it displays a valid OMB control number.

Privacy Act Statement

The Privacy Act of 1974 (P.L. 93-579) requires that you be given certain information in connection with your submission of the attached form related to a patent application or patent. Accordingly, pursuant to the requirements of the Act, please be advised that: (1) the general authority for the collection of this information is 35 U.S.C. 2(b)(2); (2) furnishing of the information solicited is voluntary; and (3) the principal purpose for which the information is used by the U.S. Patent and Trademark Office is to process and/or examine your submission related to a patent application or patent. If you do not furnish the requested information, the U.S. Patent and Trademark Office may not be able to process and/or examine your submission, which may result in termination of proceedings or abandonment of the application or expiration of the patent.

The information provided by you in this form will be subject to the following routine uses:

- 1. The information on this form will be treated confidentially to the extent allowed under the Freedom of Information Act (5 U.S.C. 552) and the Privacy Act (5 U.S.C 552a). Records from this system of records may be disclosed to the Department of Justice to determine whether disclosure of these records is required by the Freedom of Information Act.
- 2. A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement negotiations.
- 3. A record in this system of records may be disclosed, as a routine use, to a Member of Congress submitting a request involving an individual, to whom the record pertains, when the individual has requested assistance from the Member with respect to the subject matter of the record.
- 4. A record in this system of records may be disclosed, as a routine use, to a contractor of the Agency having need for the information in order to perform a contract. Recipients of information shall be required to comply with the requirements of the Privacy Act of 1974, as amended, pursuant to 5 U.S.C. 552a(m).
- 5. A record related to an International Application filed under the Patent Cooperation Treaty in this system of records may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant to the Patent Cooperation Treaty.
- 6. A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
- 7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or his/her designee, during an inspection of records conducted by GSA as part of that agency's responsibility to recommend improvements in records management practices and programs, under authority of 44 U.S.C. 2904 and 2906. Such disclosure shall be made in accordance with the GSA regulations governing inspection of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make determinations about individuals.
- 8. A record from this system of records may be disclosed, as a routine use, to the public after either publication of the application pursuant to 35 U.S.C. 122(b) or issuance of a patent pursuant to 35 U.S.C. 151. Further, a record may be disclosed, subject to the limitations of 37 CFR 1.14, as a routine use, to the public if the record was filed in an application which became abandoned or in which the proceedings were terminated and which application is referenced by either a published application, an application open to public inspection or an issued patent.
- 9. A record from this system of records may be disclosed, as a routine use, to a Federal, State, or local law enforcement agency if the USPTO becomes aware of a violation or retortial violation of law or regulation.

	Application No.	Applicant(s)	
	14/957,501	TONG ET AL	
Notice of Allowability	Examiner JOANNIE A. GARCIA	Art Unit 2895	AIA (First Inventor to File) Status No

The MAILING DATE of this communication appears on the cover sheet with the correspondence address claims being allowable, PROSECUTION ON THE MERITS IS (OR REMAINS) CLOSED in this application. If not included ewith (or previously mailed), a Notice of Allowance (PTOL-85) or other appropriate communication will be mailed in due course. THIS TICE OF ALLOWABILITY IS NOT A GRANT OF PATENT RIGHTS. This application is subject to withdrawal from issue at the initiat ne Office or upon petition by the applicant. See 37 CFR 1.313 and MPEP 1308.						
☐ This communication is responsive to <u>claims, remarks, and TD filed 02/08/2016</u> . ☐ A declaration(s)/affidavit(s) under 37 CFR 1.130(b) was/were filed on						
 An election was made by the applicant in response to a restriction recrequirement and election have been incorporated into this action. 						
3. The allowed claim(s) is/are <u>2-34</u> . As a result of the allowed claim(s), y Highway program at a participating intellectual property office for the http://www.uspto.gov/patents/init_events/pph/index.jsp or send an inq	corresponding application. For more information, please see					
4. Acknowledgment is made of a claim for foreign priority under 35 U.S.	C. § 119(a)-(d) or (f).					
Certified copies:						
a) ☐ All b) ☐ Some *c) ☐ None of the:						
 Certified copies of the priority documents have been rec 	reived.					
Certified copies of the priority documents have been rec	eived in Application No					
3. \square Copies of the certified copies of the priority documents h	nave been received in this national stage application from the					
International Bureau (PCT Rule 17.2(a)).						
* Certified copies not received:						
Applicant has THREE MONTHS FROM THE "MAILING DATE" of this connoted below. Failure to timely comply will result in ABANDONMENT of the THIS THREE-MONTH PERIOD IS NOT EXTENDABLE.						
5. CORRECTED DRAWINGS (as "replacement sheets") must be subm	itted.					
including changes required by the attached Examiner's Amenda Paper No./Mail Date	nent / Comment or in the Office action of					
Identifying indicia such as the application number (see 37 CFR 1.84(c)) sho each sheet. Replacement sheet(s) should be labeled as such in the header						
 DEPOSIT OF and/or INFORMATION about the deposit of BIOLOGIC attached Examiner's comment regarding REQUIREMENT FOR THE D 						
Attachment(a)						
Attachment(s) 1. ☐ Notice of References Cited (PTO-892)	5. Examiner's Amendment/Comment					
2. X Information Disclosure Statements (PTO/SB/08),	6. 🛮 Examiner's Statement of Reasons for Allowance					
Paper No./Mail Date <u>See Continuation Sheet</u> 3. Examiner's Comment Regarding Requirement for Deposit	7. Other					
of Biological Material						
4. Interview Summary (PTO-413), Paper No./Mail Date						
/JOHN P DULKA/						
Primary Examiner, Art Unit 2895						

U.S. Patent and Trademark Office PTOL-37 (Rev. 08-13) 20160511

Notice of Allowability

Part of Paper No./Mail Date

Continuation of Attachment(s) 2. Information Disclosure Statements (PTO/SB/08), Paper No./Mail Date: 02/16/2016, and 03/11/2016.

Application/Control Number: 14/957,501 Page 2

Art Unit: 2895

DETAILED ACTION

The present application, filed on or after March 16, 2013, is being examined under the first inventor to file provisions of the AIA.

Status of Application

In response to the Office action mailed 02/03/2016 ("02-03-16 OA"), Applicant's reply and Terminal Disclaimer filed on 02/08/2016 ("02-08-16 Reply").

Pending claims 2-34 of which claims 2, 22, 32, and 33, are independent.

Information Disclosure Statements (IDS)

The IDS filed 02/16/2016, and 03/11/2016, have been considered, and a signed and dated copy of each, is included herein.

It is noted that the prosecution history for the following related application has been reviewed and considered: 14/754,111, 14/197,070, 14/197,056, 13/341,273, 12/954,740, 12/720,368, 11/980,664, 10/913,441, 11/980,664, 11/980,415, 12/493,957, 10/762,318, 10/460,418, and 09/505,283.

Response to Arguments

- Applicant's reply and Terminal Disclaimer have overcome the nonstatutory obviousness-type double patenting rejection(s) as set forth in pages 2-20 of the 02-03-16 OA.
- Applicants' Remarks (pages 6-8) in the 02-08-16 Reply with respect the nonstatutory obviousness-type double patenting rejection(s), have been fully considered and are persuasive.

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Art Unit: 2895

Reasons for Allowance

Claims 2-34 are allowed.

The following is an examiner's statement of reasons for allowance:

Independent claim 2 is allowed, because the prior art of record, singularly or in combination, does not disclose or suggest, in combination with the other claimed elements in claim 2, terminating the planarized insulating material with a nitrogencontaining species by at least one of:

a nitrogen-containing plasma during the etching, and

a nitrogen-containing solution after the etching;

bringing a first material into direct contact with the terminated and planarized insulating material outside the reaction space; and

forming a chemical bond between the terminated and planarized insulating material and the first material.

Claims 3-21 are allowed, because they depend on allowed claim 2.

Independent claim 22 is allowed, because the prior art of record, singularly or in combination, does not disclose or suggest, in combination with the other claimed elements in claim 22, wherein the first bonding surface is an etched surface terminated with a nitrogen- containing species, and

wherein the first and second bonding surfaces are in direct contact with each other and bonded together with a chemical bond without any intervening adhesive.

Claims 23-31 are allowed, because they depend on allowed claim 22.

Independent claim 32 is allowed, because the prior art of record, singularly or in combination, does not disclose or suggest, in combination with the other claimed elements in claim 32, terminating the planarized insulating material with a nitrogencontaining species by at least one of:

a nitrogen-containing plasma during the etching, and

a nitrogen-containing solution after the etching;

bringing a first material into direct contact with the terminated and planarized insulating material; and

forming a chemical bond between the terminated and planarized insulating material and the first material.

Independent claim 33 is allowed, because the prior art of record, singularly or in combination, does not disclose or suggest, in combination with the other claimed elements in claim 33, *terminating the planarized insulating material with a species*,

bringing into direct contact with the terminated and planarized insulating material and a first material outside of said plasma chamber; and

forming a chemical bond between the terminated and planarized insulating material and the first material.

Claim 34 is allowed, because it depends on allowed claim 33.

Therefore, the prior-art of record neither anticipates nor renders obvious the claimed subject matter.

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Any comments considered necessary by applicant must be submitted no later than the payment of the issue fee and, to avoid processing delays, should preferably accompany the issue fee. Such submissions should be clearly labeled "Comments on Statement of Reasons for Allowance."

Conclusion

When responding to this office action, applicants are advised to provide the examiner with the line numbers and page numbers in the application and/or references cited to assist the examiner in locating appropriate paragraphs.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Joannie García, whose telephone number is (571) 272-1861. The examiner can normally be reached on Monday through Friday.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Mohammad K. Islam, can be reached on (571) 270-5878. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

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Art Unit: 2895

/JOANNIE A GARCIA/ Examiner, Art Unit 2895

/JOHN P DULKA/ Primary Examiner, Art Unit 2895



UNITED STATES DEPARTMENT OF COMMERCE United States Patent and Trademark Office Address: COMMISSIONER FOR PATENTS P.O. Box 1450 Alexandria, Virginia 22313-1450 www.uspto.gov

BIB DATA SHEET

CONFIRMATION NO. 1033

SERIAL NUM	IBER	FILING	_371(c)		CLASS	GR	OUP ART	UNIT	ATTO	DRNEY DOCKET
14/957,50)1	DAT 12/02/2			257		2895		T	NO. SSRA.012C1
		RUL	E							
	APPLICANTS ZIPTRONIX, INC., Morrisville, NC;									
Qin-Yi To Gaius Gi	INVENTORS Qin-Yi Tong, Durham, NC; Gaius Gillman Fountain JR., Youngsville, NC; Paul M. Enquist, Cary, NC;									
This appl wh wh wh wh wh wh	** CONTINUING DATA ***********************************									
** FOREIGN A	PPLICA	ATIONS *****	******	******	*					
** IF REQUIRE 12/11/20		EIGN FILING	LICENS	E GRA	NTED **					
Foreign Priority claim 35 USC 119(a-d) con Verified and	ditions met	Yes No Yes No A GARCIA	☐ Met at Allowa	ter ince	STATE OR COUNTRY		HEETS WINGS	TOT. CLAI	MS	INDEPENDENT CLAIMS
	Examiner's		Initials		NC		9	33	1	4
ADDRESS										
& Bear, L 2040 ma Fourteen Irvine, CA	Knobbe, Martens, Olson & Bear, LLP 2040 main Street Fourteenth Floor Irvine, CA 92614 UNITED STATES									
TITLE										
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							☐ Other			
							☐ Credit			

BIB (Rev. 05/07).

	Application No.	14/957,501
INFORMATION DISCLOSURE	Filing Date	December 2, 2015
STATEMENT BY APPLICANT	First Named Inventor	Tong et al.
STATEMENT BY AFFEIGANT	Art Unit	2895
(Multiple sheets used when necessary)	Examiner	Joannie A. Garcia
SHEET 1 OF 22	Attorney Docket No.	TSSRA.012C1

	U.S. PATENT DOCUMENTS							
Examiner Initials	Cite No.	Document Number Number - Kind Code (if known) Example: 1,234,567 B1	Publication Date MM-DD-YYYY	Name	Pages, Columns, Lines Where Relevant Passages or Relevant Figures Appear			
	1	3,423,823	01-28-1969	Ansley				
	2	3,488,834	01-13-1970	Baird				
	3	3,508,980	04-28-1970	Jackson, Jr. et al.				
	4	3,534,467	10-20-1970	Sachs et al.				
	5	3,579,391	05-18-1971	Buie				
	6	3,587,166	06-28-1971	Alexander et al.				
	7	3,602,981	09-07-1971	Kool				
	8	3,607,466	09-21-1971	Miyazaki				
	9	3,640,807	02-08-1972	Van Dijk				
	10	3,647,581	03-07-1972	Mash				
	11	3,888,708	06-10-1975	Wise et al.				
	12	4,416,054	11-22-1983	Thomas et al.				
	13	4,612,083	09-16-1986	Yasumoto et al.				
	14	4,617,160	10-14-1986	Belanger et al.				
	15	4,649,630	03-13-1987	Boland et al.				
	16	4,700,466	10-20-1987	Nakagawa et al.				
	17	4,754,544	07-05-1988	Hanak				
	18	4,829,018	05-09-1989	Wahlstrom				
	19	4,935,386	06-19-1990	Nakagawa				
	20	4,939,101	07-03-1990	Black et al.				
	21	4,962,879	10-16-1990	Gösele et al.				
	22	4,963,505	10-16-1990	Fujii et al.				
	23	4,970,175	11-13-1990	Haisma et al.				
	24	4,978,421	12-18-1990	Bassous et al.				
	25	4,983,251	01-08-1991	Haisma et al.				
	26	4,992,149	02-12-1991	Nguyen				
	27	5,007,071	04-09-1991	Nakano				
	28	5,013,471	05-07-1991	Ogawa				
	29	5,024,723	06-18-1991	Gösele et al.				

Examiner Signature	/Joannie Garcia/	Date Considered 05/11/2016
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^{*}Examiner: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

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STATEMENT BY APPLICANT	Art Unit	2895
(Multiple sheets used when necessary)	Examiner	Joannie A. Garcia
SHEET 2 OF 22	Attorney Docket No.	TSSRA.012C1

U.S. PATENT DOCUMENTS					
Examiner Initials	Cite No.	Document Number Number - Kind Code (if known) Example: 1,234,567 B1	Publication Date MM-DD-YYYY	Name	Pages, Columns, Lines Where Relevant Passages or Relevant Figures Appear
	30	5,034,343	07-23-1991	Rouse et al.	
	31	5,070,026	12-03-1991	Greenwald et al.	
	32	5,071,792	12-10-1991	Van Vonno et al.	
	33	5,081,061	01-14-1992	Rouse et al.	
	34	5,087,585	02-11-1992	Hayashi	
	35	5,089,431	02-18-1992	Slatter et al.	
	36	5,121,706	06-16-1992	Nichols et al.	
	37	5,162,251	11-10-1992	Poole et al.	
	38	5,183,783	02-02-1993	Ohta et al.	
	39	5,196,375	03-23-1993	Hoshi	
	40	5,207,864	05-04-1993	Bhat et al.	
	41	5,234,860	08-10-1993	Gluck	
	42	5,236,118	08-17-1993	Bower et al.	
	43	5,238,875	08-24-1993	Ogino	
	44	5,266,511	11-30-1993	Takao	
	45	5,266,824	11-30-1993	Abe	
	46	5,270,259	12-14-1993	Ito et al.	
	47	5,270,261	12-14-1993	Bertin et al.	
	48	5,272,104	12-21-1993	Schrantz et al.	
	49	5,284,707	02-08-1994	Ogawa et al.	
	50	5,321,301	06-14-1994	Sato et al.	
	51	5,324,566	06-28-1994	Ogawa et al.	
	52	5,324,687	06-28-1994	Wojnarowski	
	53	5,354,695	10-11-1994	Leedy	
	54	5,362,659	11-08-1994	Cartagena	
	55	5,376,579	12-27-1994	Annamalai	
	56	5,380,681	01-10-1995	Hsu	
	57	5,383,993	01-24-1995	Katada et al.	
	58	5,407,856	04-18-1995	Quenzer et al.	

Examiner Signature /Joannie Garcia/	Date Considered	05/11/2016
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STATEMENT BY AFFLICANT	Art Unit	2895
(Multiple sheets used when necessary)	Examiner	Joannie A. Garcia
SHEET 3 OF 22	Attorney Docket No.	TSSRA.012C1

U.S. PATENT DOCUMENTS					
Examiner Initials	Cite No.	Document Number Number - Kind Code (if known) Example: 1,234,567 B1	Publication Date MM-DD-YYYY	Name	Pages, Columns, Lines Where Relevant Passages or Relevant Figures Appear
	59	5,413,955	05-09-1995	Lee et al.	
	60	5,420,449	05-30-1995	Oji	
	61	5,421,953	06-06-1995	Nagakubo et al.	
	62	5,422,302	06-06-1995	Yonehara	
	63	5,427,638	06-27-1995	Goetz et al.	
	64	5,432,729	07-11-1995	Carson et al.	
	65	5,437,894	08-01-1995	Ogawa et al.	
	66	5,441,591	08-15-1995	Imthum et al.	
	67	5,441,911	08-15-1995	Malhi	
	68	5,459,104	10-17-1995	Sakai	
	69	5,460,659	10-24-1995	Krut	
	70	5,485,540	01-16-1996	Eda	
	71	5,489,554	02-06-1996	Gates	
	72	5,497,033	03-05-1996	Fillion et al.	
	73	5,503,704	04-02-1996	Bower et al.	
	74	5,514,235	05-07-1996	Mitani et al.	
	75	5,516,727	05-14-1996	Broom	
	76	5,517,754	05-21-1996	Bellstein, Jr., et. al.	
	77	5,523,602	06-04-1996	Horiuchi	
	78	5,534,465	07-09-1996	Frye et al.	
	79	5,543,648	08-06-1996	Miyawaki	
	80	5,546,494	08-13-1996	Eda	
	81	5,547,896	08-20-1996	Linn et al.	
	82	5,548,178	08-20-1996	Eda et al.	
	83	5,561,303	10-01-1996	Schrantz et al.	
	84	5,563,084	10-08-1996	Ramm et al.	
	85	5,567,657	10-22-1996	Wonjnarowski et al.	
	86	5,569,620	10-29-1996	Linn et al.	
	87	5,580,407	12-03-1996	Haisma et al.	

Examiner Signature /Joannie Garcia/	Date Considered 05/11/	2016
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	Application No.	14/957,501	
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STATEMENT BY AFFEIGANT	Art Unit	2895	
(Multiple sheets used when necessary)	Examiner	Joannie A. Garcia	
SHEET 4 OF 22	Attorney Docket No.	TSSRA.012C1	

U.S. PATENT DOCUMENTS					
Examiner Initials	Cite No.	Document Number Number - Kind Code (if known) Example: 1,234,567 B1	Publication Date MM-DD-YYYY	Name	Pages, Columns, Lines Where Relevant Passages or Relevant Figures Appear
	88	5,591,678	01-07-1997	Bendik et al.	
	89	5,627,106	05-06-1997	Hsu	
	90	5,647,932	07-15-1997	Taguchi et al.	
	91	5,650,353	07-22-1997	Yoshizawa et al.	
	92	5,652,436	07-29-1997	Stoner et al.	
	93	5,653,019	08-05-1997	Bernhardt et al.	
	94	5,659,192	08-19-1997	Sarma et al.	
	95	5,661,316	08-26-1997	Kish, Jr., et al.	
	96	5,661,901	09-02-1997	King	
	97	5,666,706	09-16-1997	Tomita et al.	
	98	5,668,057	09-16-1997	Eda et al.	
	99	5,672,240	09-30-1997	Stoner et al.	
	100	5,673,478	10-07-1997	Beene et al.	
	101	5,698,471	12-16-1997	Namba et al.	
	102	5,741,733	04-21-1998	Bertagnolli et al.	
	103	5,747,857	05-05-1998	Eda et al.	
	104	5,753,529	05-19-1998	Chang et al.	
	105	5,755,914	05-26-1998	Yonehara	
	106	5,759,753	06-02-1998	Namba et al.	
	107	5,760,478	06-02-1998	Bazso et al.	
	108	5,763,318	06-09-1998	Bertin et al.	
	109	5,766,984	06-16-1998	Ramm et al.	
	110	5,771,555	06-30-1998	Eda et al.	
	111	5,783,477	07-21-1998	Kish, Jr. et al.	
	112	5,785,874	07-28-1998	Eda et al.	
	113	5,793,115	08-11-1998	Zavracky et al.	
	114	5,804,086	09-08-1998	Bruel	
	115	5,807,783	09-15-1998	Gaul et al.	
	116	5,821,665	10-13-1998	Onishi et al.	

Examiner Signature	Joannie Garcia/	Date Considered	05/11/2016

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PTO/SB/08 Equivalent

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STATEMENT BY APPLICANT	First Named Inventor	Tong et al.
STATEMENT BY ALL LICANT	Art Unit	2895
(Multiple sheets used when necessary)	Examiner	Joannie A. Garcia
SHEET 5 OF 22	Attorney Docket No.	TSSRA.012C1

	Francisco	0:4-	Document Number	Publication		Pages, Columns, Lines Where
	Examiner Initials	Cite No.	Number - Kind Code (if known) Example: 1,234,567 B1	Date MM-DD-YYYY	Name	Relevant Passages or Relevant Figures Appear
		117	5,841,197	11-24-1998	Adamic, Jr.	
		118	5,849,627	12-15-1998	Linn et al.	
		119	5,851,894	12-22-1998	Ramm	
		120	5,858,876	01-12-1999	Chew	
		121	5,866,942	02-02-1999	Suzuki et al.	
		122	5,869,354	02-09-1999	Leedy et al.	
		123	5,872,025	02-16-1999	Cronin et al.	
		124	5,876,497	03-02-1999	Atoji	
		125	5,877,034	03-02-1999	Ramm et al.	
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		127	5,877,516	03-02-1999	Mermagen et al.	
		128	5,880,010	03-09-1999	Davidson	
		129	5,889,302	03-30-1999	Liu	
hang		130	5,902,118	05-11-1999	Hubner	
pplied		131	5,903,018	05-11-1999	Shimawaki	
ocum	eni	132	5,904,860	05-18-1999	Nagakubo et al.	
J.G./		1335	910699 -5,810,660	06-08-1999	Namba et al.	
5/11/20	រា6	134	5,915,167	06-22-1999	Leedy	
		135	5,915,193	06-22-1999	Tong et al	
		136	5,920,142	07-06-1999	Onishi et al.	
		137	5,936,280	08-10-1999	Liu	
		138	5,966,622	10-12-1999	Levine et al.	
		139	5,980,770	11-09-1999	Ramachandran et al.	
		140	5,982,010	11-09-1999	Namba et al.	
		141	5,990,472	11-23-1999	Rinne	
		142	5,990,562	11-23-1999	Vallett	
		143	5,991,989	11-30-1999	Onishi et al.	
		144	6,004,865	12-21-1999	Horiuchi	
		145	6,004,866	12-21-1999	Nakano et al.	

Examiner Signature Date Considered 05/11/2016 /Joannie Garcia/

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PTO/SB/08 Equivalent

	Application No.	14/957,501
INFORMATION DISCLOSURE	Filing Date	December 2, 2015
STATEMENT BY APPLICANT	First Named Inventor	Tong et al.
STATEMENT BY APPLICANT	Art Unit	2895
(Multiple sheets used when necessary)	Examiner	Joannie A. Garcia
SHEET 6 OF 22	Attorney Docket No.	TSSRA.012C1

				U.S. PATENT	TENT DOCUMENTS		
	Examiner Initials	Cite No.	Document Number Number - Kind Code (if known) Example: 1,234,567 B1	Publication Date MM-DD-YYYY	Name	Pages, Columns, Lines Where Relevant Passages or Relevant Figures Appear	
Chang	nes eer	146	6,018,211	01-25-2000	Kanaboshi et al.		
applie	="	147	6,048,752	04-11-2000	Lindemann		
docun		148	6,054,371 04/2000	09-29-1998	Tsuchiaki et al.		
/J.G./		149	6,071,761	06-06-2000	Jacobs		
05/11/2)16	150	6,087,719 07/2000	.04.23.4008	Tsunashima		
		151	6,087,760	07-11-2000	Yamaguchi et al.		
		152	6,103,009	08-15-2000	Atoji		
		153	6,120,917	09-19-2000	Eda		
		154	6,133,640	10-17-2000	Leedy		
		155	6,136,691 10/2000	-05-26-4008	Chen		
		156	6,143,628	11-07-2000	Sato et al.		
		157	6,146,992	11-14-2000	Lauterbach et al.		
		158	6,153,445	11-28-2000	Yamazaki et al.		
		159	6,153,495	11-28-2000	Kub et al.		
		160	6,154,940	12-05-2000	Onishi et al.		
		161	6,156,624	12-05-2000	Yamagata et al.		
		162	6,165,817	12-26-2000	Akram et al.		
		163	6,180,496	01-30-2001	Farrens et al.		
		164	6,190,778	02-20-2001	Batz-Sohn et al.		
		165	6,194,323	02-27-2001	Downey et al.		
		166	6,197,663	03-06-2001	Chandross et al.		
		167	6,198,159	03-06-2001	Hosoma et al.		
		168	6,236,141	05-22-2001	Sato et al.		
		169	6,242,324	06-05-2001	Kub		
		170	6,246,068	06-12-2001	Sato et al.		
		171	6,255,731	07-03-2001	Ohmi et al.		
		172	6,255,899	07-03-2001	Bertin et al.		
		173	6,270,202	08-07-2001	Namba et al.		
		174	6,274,892	08-14-2001	Kub et al.		

Examiner Signature /Joannie Garcia/ Date Considered 05/11/2016

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PTO/SB/08 Equivalent

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STATEMENT BY APPLICANT	Art Unit	2895
(Multiple sheets used when necessary)	Examiner	Joannie A. Garcia
SHEET 7 OF 22	Attorney Docket No.	TSSRA.012C1

	U.S. PATENT DOCUMENTS							
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nges	175	6,284,085	09-04-2001	Gwo				
ed to	176	6,313,012	11-06-2001	Horiuchi				
ment	177	6,316,332	11-13-2001	Lo et al.				
/	178	6,323,108	11-27-2001	Kub et al.				
016	179	6,328,796	12-11-2001	Kub				
	180	6,328,841 12/2001	~%9=24=+997	Klumpp et al.				
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STATEMENT BY AFFEIGANT	Art Unit	2895
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Examiner Signature /Joannie Garcia/ Date Considered 05/11/2016
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^{*}Examiner: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

	Application No.	14/957,501
INFORMATION DISCLOSURE	Filing Date	December 2, 2015
STATEMENT BY APPLICANT	First Named Inventor	Tong et al.
STATEMENT BY AFFEIGANT	Art Unit	2895
(Multiple sheets used when necessary)	Examiner	Joannie A. Garcia
SHEET 21 OF 22	Attorney Docket No.	TSSRA.012C1

		NON PATENT LITERATURE DOCUMENTS						
Examiner Initials	Litam (hook magazina jaurnal carial cumpacium actalag ata) data paga(c) valuma jacua numbar(c) L							
	440	U. Gösele et al., History and Future of Semiconductor Wafer Bonding, 47-48 Solid State Phenomena 33 (1996).						
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Examiner Signature	/Joannie Garcia/	Date Considered	05/11/2016
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^{*}Examiner: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

PTO/SB/08 Equivalent

	Application No.	14/957,501
INFORMATION DISCLOSURE	Filing Date	December 2, 2015
STATEMENT BY APPLICANT	First Named Inventor	Tong et al.
STATEMENT BY APPLICANT	Art Unit	2895
(Multiple sheets used when necessary)	Examiner	Joannie A. Garcia
SHEET 22 OF 22	Attorney Docket No.	TSSRA.012C1

		NON PATENT LITERATURE DOCUMENTS					
Examiner Initials	itom (hook magazina journal carial symposium catalog ata) data paga(s) valuma issua numbar(s)						
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	460	Stefan Bengtsson et al., Journal of Electronic Materials, vol. 29, No. 7, "Room Temperature Wafer Bonding Of Silicon Oxidized Silicon, and Crystalline Quartz," 2000, pp. 909-915.					
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PTO/SB/08 Equivalent

	Application No.	14/957,501
INFORMATION DISCLOSURE	Filing Date	December 2, 2015
STATEMENT BY APPLICANT	First Named Inventor	Tong et al.
STATEMENT BY AFFEIGAINT	Art Unit	2895
(Multiple sheets used when necessary)	Examiner	Joannie A. Garcia
SHEET 1 OF 1	Attorney Docket No.	TSSRA.012C1

	U.S. PATENT DOCUMENTS						
Examiner Initials	Cite No.	Document Number Number - Kind Code (if known) Example: 1,234,567 B1	Publication Date MM-DD-YYYY	Name	Pages, Columns, Lines Where Relevant Passages or Relevant Figures Appear		

	FOREIGN PATENT DOCUMENTS						
Examiner Initials	Cite No.	Foreign Patent Document Country Code-Number-Kind Code Example: JP 1234567 A1	Publication Date MM-DD-YYYY	Name	Pages, Columns, Lines Where Relevant Passages or Relevant Figures Appear	T¹	

		NON PATENT LITERATURE DOCUMENTS						
Examiner Initials	item (hook magazine journal serial symposium estalog etc.) date page(s) volume issue number(s)							
	1	AMIRFEIZ et al., "Formation of silicon structures by plasma activated wafer bonding," Vol. 99-2, Abstract No. 963, 1999 Joint Int'l Meeting of the Electrochecm. Soc'y, December 16, 1999.						
	2	AMIRFEIZ et al., "Formation of silicon structures by plasma activated wafer bonding," Electrochemical Society Proceedings: Semiconductor Wafer Bonding: Science, Technology, and Applications V, Volume 99-35, (1999), pp. 29-39.						
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Examiner Signature /Joannie Garcia/ Date Considered 05/11/2016

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TONG ET AL.

Applicant(s)/Patent Under Reexamination

Examiner

Art Unit

JOANNIE A GARCIA

2895

CPC						
Symbol					Туре	Version
H01L	29	7	16		F	2013-01-01
Y10T	156	9	1043		А	2015-01-15
Y10T	156	9	10		А	2015-01-15
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H01L	21		31105		1	2013-01-01
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H01L	24	1	83		ı	2013-01-01
H01L	2224	7	8319		А	2013-01-01
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H01L	2224	7	83894		А	2013-01-01
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/JOANNIE A GARCIA/ Examiner.Art Unit 2895				05/11/2016	Total Claims Allowed:	
(Assistant Examiner)				(Date)	3	3
/JOHN P [Primary E	DULKA/ xaminer.Art Unit 28	95		05/13/2016	O.G. Print Claim(s)	O.G. Print Figure
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TONG ET AL.

Applicant(s)/Patent Under Reexamination

Examiner

JOANNIE A GARCIA

Art Unit

2895

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H01L	2924	1	0132	Α	2013-01-01
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CPC Com	bination Sets					
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/JOANNIE A GARCIA/ Examiner.Art Unit 2895	05/11/2016	Total Claims Allowed:				
(Assistant Examiner)	(Date)	33				
/JOHN P DULKA/ Primary Examiner.Art Unit 2895	05/13/2016	O.G. Print Claim(s)	O.G. Print Figure			
(Primary Examiner)	(Date)	1	4A-4E			

U.S. Patent and Trademark Office

Application/Control No.	Applicant(s)/Patent Under Reexamination
14957501	TONG ET AL.
Examiner	Art Unit

2895

	US ORIGINAL CLASSIFICATION						INTERNATIONAL CLASSIFICATION									
	CLASS			SUBCLASS					С	LAIMED		NON-CLAIMED				
						Н	0	1	L	21 / 30 (2006.01.01)						
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/JOANNIE A GARCIA/ Examiner.Art Unit 2895	05/11/2016	Total Claims Allowed:			
(Assistant Examiner)	(Date)	33			
/JOHN P DULKA/ Primary Examiner.Art Unit 2895	05/13/2016	O.G. Print Claim(s)	O.G. Print Figure		
(Primary Examiner)	(Date)	1	4A-4E		

U.S. Patent and Trademark Office



Application/Control No. 14957501 Examiner	Applicant(s)/Patent Under Reexamination
14957501	TONG ET AL.
Examiner	Art Unit
JOANNIE A GARCIA	2895

☐ Claims renumbered in the same order as presented by applicant ☐ CPA ☒ T.D. ☐ R.1.47															
Final	Original	Final	Original	Final	Original	Final	Original	Final	Original	Final	Original	Final	Original	Final	Original
	1	16	17	32	33										
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/JOANNIE A GARCIA/ Examiner.Art Unit 2895	05/11/2016	Total Claims Allowed:			
(Assistant Examiner)	(Date)	33			
/JOHN P DULKA/ Primary Examiner.Art Unit 2895	05/13/2016	O.G. Print Claim(s)	O.G. Print Figure		
(Primary Examiner)	(Date)	1	4A-4E		

U.S. Patent and Trademark Office

EAST Search History

EAST Search History (Prior Art)

Ref #	Hits	Search Query	DBs	Default Operator	Plurals	Time Stamp
L1	480415	(bond\$).clm.	US- PGPUB; USPAT; USOCR		OFF	2016/05/11 20:05
L2	59338	(anneal\$).clm.	US- PGPUB; USPAT; USOCR		OFF	2016/05/11 20:05
L3	880726	(heat or heats or heated or heating).clm.	US- PGPUB; USPAT; USOCR		OFF	2016/05/11 20:05
L4	143	L1 with L2 with L3	US- PGPUB; USPAT; USOCR	OR	OFF	2016/05/11 20:05
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S21	56069	(anneal\$).clm.	US- PGPUB; USPAT; USOCR	OR	OFF	2015/07/22 15:54
S22	839493	(heat or heats or heated or heating).clm.	US- PGPUB; USPAT; USOCR	OR	OFF	2015/07/22 15:54
S23	138	S20 with S21 with S22	US- PGPUB; USPAT; USOCR	OR	OFF	2015/07/22 15:55
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S32	1	("20150303263").PN.	US- PGPUB; USPAT; USOCR	OR	OFF	2015/11/10 16:41
S33	1	("5451547").PN.	US- PGPUB; USPAT; USOCR	OR	OFF	2015/11/10 16:41
S34	1286659	adhesive	US- PGPUB; USPAT; USOCR	OR	ON	2015/11/10 16:41
S35	0	S33 and S34	US- PGPUB; USPAT; USOCR		ON	2015/11/10 16:41
S36	2043	terminated with species	US- PGPUB; USPAT; USOCR	OR	ON	2015/11/10 16:45
S37	1	S32 and S36	US- PGPUB; USPAT; USOCR	OR	ON	2015/11/10 16:45
S38	909961	etch\$	US- PGPUB; USPAT; USOCR	OR	ON	2015/11/10 16:48
S39	1	S33 and S38	US- PGPUB; USPAT; USOCR	OR	ON	2015/11/10 16:49
S40	5	(bond or bonds or bonded or bonding) with (etch or etches or etched or etching) with (("25" or "room temperature" or "room-temperature" or "room T" or "room-T") and ("chemical bond" or "chemical-bond"))	PGPUB; USPAT;	OR	OFF	2015/11/10 16:58
S41	28	(bond or bonds or bonded or bonding) with (etch or etches or etched or etching) with (("25" or "room temperature" or "room-temperature" or "room T" or "room-T") and ("chemical bond" or "chemical-bond"))	PGPUB; USPAT;	OR	ON	2015/11/10 16:58
S42	0	S33 and S41	US- PGPUB; USPAT; USOCR	OR	ON	2015/11/10 16:58
S43	48227	(bond or bonds or bonded or bonding) with (etch or etches or etched or etching)	US- PGPUB;	OR	ON	2015/11/10 16:58

			USPAT; USOCR			
S44	0	S33 and S43	US- PGPUB; USPAT; USOCR	OR	ON	2015/11/10 16:59
S45	2042101	bond or bonds or bonded or bonding	US- PGPUB; USPAT; USOCR	OR	ON	2015/11/10 16:59
S46	1	S33 and S45	US- PGPUB; USPAT; USOCR	OR	ON	2015/11/10 16:59
S47	567781	CMP or polish\$ or planarize or planarizes or planarized or planarizing or planarization	US- PGPUB; USPAT; USOCR	OR	ON	2015/11/10 17:04
S48	1	S33 and S47	US- PGPUB; USPAT; USOCR	OR	ON	2015/11/10 17:04
S49	3009473	plasma or gas	US- PGPUB; USPAT; USOCR	OR	ON	2015/11/10 17:09
S50	1	S33 and S49	US- PGPUB; USPAT; USOCR	OR	ON	2015/11/10 17:09
S51	1	S39 and S46 and S48 and S50	US- PGPUB; USPAT; USOCR	OR	ON	2015/11/10 17:10
S52	913511	etch\$ or VSE	US- PGPUB; USPAT; USOCR	OR	ON	2015/11/10 17:49
S53	1	S33 and S52	US- PGPUB; USPAT; USOCR	OR	ON	2015/11/10 17:49
S54	790750	specie	US- PGPUB; USPAT; USOCR	OR	ON	2015/11/10 17:53
S55	0	S33 and S54	US- PGPUB; USPAT; USOCR		ON	2015/11/10 17:54
S56	1	("5451547").PN.	US- PGPUB; USPAT; USOCR	OR	OFF	2015/11/14 10:09
S57	1	("20150303263").PN.	US- PGPUB; USPAT; USOCR	OR	OFF	2015/11/14 10:38
					<u></u>	

S58	85	"planarized insulating material"	US- PGPUB; USPAT; USOCR	OR	ON	2015/11/14 10:43
S59	1	S57 and S58	US- PGPUB; USPAT; USOCR	OR	ON	2015/11/14 10:43
S60	472066	(bond\$).clm.	US- PGPUB; USPAT; USOCR	OR	OFF	2016/01/25 18:29
S61	58187	(anneal\$).clm.	US- PGPUB; USPAT; USOCR	OR	OFF	2016/01/25 18:29
S62	865929	(heat or heats or heated or heating).clm.	US- PGPUB; USPAT; USOCR	OR	OFF	2016/01/25 18:29
S63	140	S60 with S61 with S62	US- PGPUB; USPAT; USOCR	OR	OFF	2016/01/25 18:29
S64	22	S63 AND ((H01L21/324 OR H01L21/2022 OR H01L21/26506 OR H01L21/187 OR H01L21/2007 OR H01L21/76251 OR H01L23/38 OR H01L21/02032 OR H01L21/3065 OR H01L21/3226 OR H01L21/67092).CPC.)	US- PGPUB; USPAT; USOCR	OR	OFF	2016/01/25 18:29
S65	19956	((257/459) or (257/676) or (257/684) or (257/701) or (257/702) or (257/e21.088) or (257/e21.122) or (257/e21.215) or (257/e21.218) or (257/e21.219) or (257/e21.224) or (257/e21.224) or (257/e21.243) or (257/e21.245) or (257/e21.246) or (257/e21.249) or (257/e21.33) or (257/e21.567) or (257/e23.02) or (257/e23.04)).CCLS.	US- PGPUB; USPAT; USOCR	OR	OFF	2016/01/25 18:45
S66	5551	((438/405) or (438/455) or (438/459) or (438/974) or (148/DIG12)).CCLS.	US- PGPUB; USPAT; USOCR	OR	OFF	2016/01/25 18:51
S67	24326	S65 or S66	US- PGPUB; USPAT; USOCR	OR	OFF	2016/01/25 18:51
S68	1406	(bond or bonds or bonded or bonding) with (etch or etches or etched or etching) with ("25" or "room temperature" or "room- temperature" or "room T" or "room-T")	US- PGPUB; USPAT; USOCR	OR	OFF	2016/01/25 18:52
S69	28	(bond or bonds or bonded or bonding) with (etch or etches or etched or etching) with (("25" or "room temperature" or "room- temperature" or "room T" or "room-T") and ("chemical bond" or "chemical-bond"))	PGPUB; USPAT;	• •	ON	2016/01/25 18:53
S70	4	(("4700466") or ("4983251") or ("5427638") or ("5451547")).PN.	US- PGPUB; USPAT; USOCR	OR	OFF	2016/01/25 18:58

S71	1	("20150303263").PN.	US- PGPUB; USPAT; USOCR		OFF	2016/01/25 19:30
S72	86	"planarized insulating material"	US- PGPUB; USPAT; USOCR		ON	2016/01/25 19:30
S73	1	S71 and S72	US- PGPUB; USPAT; USOCR		ON	2016/01/25 19:30
S74	1352309	nitrogen	US- PGPUB; USPAT; USOCR		ON	2016/01/25 20:26
S75	0	S70 and S74	US- PGPUB; USPAT; USOCR		ON	2016/01/25 20:26
S76	1304100	adhesive	US- PGPUB; USPAT; USOCR		ON	2016/01/25 20:27
S77	1	S70 and S76	US- PGPUB; USPAT; USOCR		ON	2016/01/25 20:27
S78	1	("8153505").PN.	US- PGPUB; USPAT; USOCR	OR	OFF	2016/01/25 21:14

EAST Search History (Interference)

Ref #	Hits	Search Query	DBs	Default Operator	Plurals	Time Stamp
L6	8028	((H01L21/324 OR H01L21/2022 OR H01L21/26506 OR H01L21/187 OR H01L21/2007 OR H01L21/76251 OR H01L23/38 OR H01L21/02032 OR H01L21/3065 OR H01L21/3226 OR H01L21/67092).CPC.)	USPAT	OR	OFF	2016/05/11 20:05
L7	12	L6 AND (bond\$ with anneal\$ with (heat or heats or heated or heating)).CLM.	USPAT	OR	OFF	2016/05/11 20:09
L9	5	(("4700466") or ("4983251") or ("5427638") or ("5451547") or ("4992149")).PN.	US- PGPUB; USPAT	OR	OFF	2016/05/11 20:09

5/11/2016 8:10:29 PM

 $\textbf{C:} \ \textbf{Users} \ \textbf{jgarcia1} \ \textbf{Documents} \ \textbf{EAST} \ \textbf{Workspaces} \ \textbf{14957501.wsp}$

Search Notes

Application/Control No.	Applicant(s)/Patent Under Reexamination
14957501	TONG ET AL.
Examiner	Art Unit
JOANNIE A GARCIA	2895

CPC- SEARCHED		
Symbol	Date	Exa mine r
H01L: 21/02032,21/185,21/187,21/2007,21/2022,21/26506,21/30,21/302,21/306,21/3065,21/3226,21/324,21/38,21/46,21/461,21/67092,21/76251,21/76254	05/11/ 2016	JAG

CPC COMBINATION SETS - SEARCHED				
Symbol	Date	Examiner		

US CLASSIFICATION SEARCHED					
Class	Subclass	Date	Examiner		
438	405,455,459,974,FOR103,FOR127,FOR388,FOR395	05/11/2016	JAG		
257	459,676,684,701,702,E21.088,E21.122,E21.215,E21.218 ,E21.219,E21.224,E21.243,E21.245,E21.249,E21.33,E23 .02,E23.04	05/11/2016	JAG		
148	DIG12	05/11/2016	JAG		

SEARCH NOTES		
Search Notes	Date	Examiner
EAST search attached	05/11/2016	JAG
Consulted Primary Examiner John P. Dulka	05/11/2016	JAG

INTERFERENCE SEARCH					
US Class/ CPC Symbol	US Subclass / CPC Group	Date	Examiner		
H01L	21/02032,21/185,21/187,21/2007,21/2022,21/26506,21/30,21/302,21/306,21/3065,21/3226,21/324,21/38,21/46, 21/461,21/67092,21/76251,21/76254	05/11/2016	JAG		

/JOANNIE A GARCIA/	
Examiner, Art Unit 2895	

PART B - FEE(S) TRANSMITTAL

Complete and send this form, together with applicable fee(s), to: Mail Mail Stop ISSUE FEE

Mail Stop ISSUE FEE Commissioner for Patents P.O. Box 1450 Alexandria, Virginia 22313-1450

Alexandria, Virginia 2 or <u>Fax</u> (571)-273-2885

INSTRUCTIONS: This form should be used for transmitting the ISSUE FEE and PUBLICATION FEE (if required). Blocks 1 through 5 should be completed where appropriate. All further correspondence including the Patent, advance orders and notification of maintenance fees will be mailed to the current correspondence address as indicated unless corrected below or directed otherwise in Block 1, by (a) specifying a new correspondence address; and/or (b) indicating a separate "FEE ADDRESS" for maintenance fee notifications.

indicated unless correct maintenance fee notifica		nerwise in Block 1, by (a					rate "FEE ADDRESS" for
CURRENT CORRESPOND	ENCE ADDRESS (Note: Use Bl	pape	e: A certificate of s) Transmittal. Things: Each additional its own certificate	l paper, su	ich as an assignmer	domestic mailings of the or any other accompanying at or formal drawing, must	
135980 7590 05/24/2016 Knobbe, Martens, Olson & Bear, LLP 2040 main Street			I he State addr trans	Cer reby certify that th es Postal Service w essed to the Mail smitted to the USP	tificate of is Fee(s) T vith sufficion Stop ISS TO (571) 2	Mailing or Transn Transmittal is being ent postage for first UE FEE address 273-2885, on the dat	nission deposited with the United t class mail in an envelope above, or being facsimile te indicated below.
Fourteenth Floo							(Depositor's name)
Irvine, CA 9261	4						(Signature)
			L				(Date)
APPLICATION NO.	FILING DATE		FIRST NAMED INVENTOR	NTOR		EY DOCKET NO.	CONFIRMATION NO.
14/957,501	12/02/2015		Qin-Yi Tong		TSS	SRA.012C1	1033
TITLE OF INVENTION	: METHOD FOR LOW	TEMPERATURE BOND	DING AND BONDED STR	UCTURE			
APPLN. TYPE	ENTITY STATUS	ISSUE FEE DUE	PUBLICATION FEE DUE	PREV. PAID ISSU	E FEE T	TOTAL FEE(S) DUE	DATE DUE
nonprovisional	UNDISCOUNTED	\$960	\$0	\$0		\$960	08/24/2016
EXAM	IINER	ART UNIT	CLASS-SUBCLASS				
GARCIA, J	OANNIE A	2895	257-684000	l			
1. Change of correspond CFR 1.363).	ence address or indication	n of "Fee Address" (37	2. For printing on the p	atent front page, lis	st	. Knobb	e, Martens,
_ ′	ondence address (or Cha	nge of Correspondence	(1) The names of up to or agents OR, alternative		nt attorneys	,	
Change of correspondence address (or Change of Correspondence Address form PTO/SB/122) attached.		(2) The name of a single firm (having as a member a registered attorney or agent) and the names of up to					
☐ "Fee Address" indication (or "Fee Address" Indication form PTO/SB/47; Rev 03-02 or more recent) attached. Use of a Customer Number is required.			2 registered patent attor listed, no name will be	neys or agents. If printed.	no name is	3	
			THE PATENT (print or typ	· *		.6. 11 1 .1 1	
recordation as set fort	th in 37 CFR 3.11. Comp	pletion of this form is NO	T a substitute for filing an	itent. If an assign assignment.	ee is ident	illed below, the do	cument has been filed for
(A) NAME OF ASSI	GNEE		(B) RESIDENCE: (CITY	and STATE OR C	COUNTRY	")	
ZIPTRONIX, INC.			Morrisville, NC				
Please check the appropr	riate assignee category or	categories (will not be pr	rinted on the patent): \Box	Individual Co	orporation o	or other private gro	up entity 🖵 Government
4a. The following fee(s)	are submitted:	41	o. Payment of Fee(s): (Plea	se first reapply ar	ny previou	ısly paid issue fee s	hown above)
Issue Fee	No small entity discount p	normittad)	A check is enclosed.	L Form DTO 2029	is ottochou	A	
	of Copies		Payment by credit card. Form PTO-2038 is attached. The director is hereby authorized to charge the required fcc(s), any deficiency, or credits any overpayment, to Deposit Account Number				
			overpayment, to Depo	sit Account Numbe	er <u>11-1</u> 4	410 (enclose an	extra copy of this form).
	tus (from status indicated						
Applicant certifying	ng micro entity status. Se	e 37 CFR 1.29	NOTE: Absent a valid cere fee payment in the micro	tification of Micro	Entity Sta	itus (see forms PTO epted at the risk of	/SB/15A and 15B), issue application abandonment.
☐ Applicant asserting small entity status. See 37 CFR 1.27		NOTE: If the application was previously under micro entity status, checking this box will be taken to be a notification of loss of entitlement to micro entity status.					
Applicant changing	ng to regular undiscounted	d fee status.	NOTE: Checking this box entity status, as applicable	will be taken to b		•	lement to small or micro
NOTE: This form must b	oe signed in accordance v	vith 37 CFR 1.31 and 1.33	3. See 37 CFR 1.4 for signa	ture requirements	and certific	cations.	
Authorized Signature	/Paul Stellma	an/		Date	May 25	, 2016	
Typed or printed name	Paul	Stellman		Registration N	Jo	68,287	

Page 320 of 331

Page 2 of 3
OMB 0651-0033

SAMSUNG EXHIBIT 1002

PTOL-85 Part B (10-13) Approved for use through 10/31/2013.

U.S. Patent and Traden**Samsungs.Welfn.wenssas: Bondling**E

TSSRA.012C1 PATENT

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Inventor : Tong et al.

App. No. : 14/957,501

Filed : December 2, 2015

For : METHOD FOR LOW

TEMPERATURE BONDING AND

BONDED STRUCTURE

Examiner : Joannie A. Garcia

Art Unit : 2895

Conf. No. : 1033

COMMENTS ON REASONS FOR ALLOWANCE

Mail Stop AF

Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450

Dear Sir:

Applicants note that the Reasons for Allowance stated in the Notice of Allowance mailed May 24, 2016, pursuant to MPEP 1302.14, are not intended to necessarily state all the reasons for allowance or all the details why the claims are allowed, and therefore does not specifically or impliedly state that all the reasons for allowance are set forth. Furthermore, the fact that Applicants have not specifically commented on the Examiner's characterizations of the reasons for allowance in the Statement of Reasons for Allowance shall not be taken as an acknowledgement of the accuracy of such characterizations. Accordingly, Applicants submit that the allowed claims may be allowable for reasons other than those stated in the Statement for Reasons for Allowance and that Applicants are not bound by the reasons for allowance provided therein.

Application No.: 14/957,501

Filing Date: December 2, 2015

Please charge any additional fees, including any fees for additional extension of time, or credit overpayment to Deposit Account No. 11-1410.

Respectfully submitted,

KNOBBE, MARTENS, OLSON & BEAR, LLP

Dated: May 25, 2016 By: /Paul Stellman/

Paul Stellman Registration No. 68,287 Attorney of Record Customer No. 20,995 (415) 954-4114

23409915 052516

Electronic Patent Application Fee Transmittal						
Application Number:	14957501					
Filing Date:	02-	Dec-2015				
Title of Invention:	METHOD FOR LOW TEMPERATURE BONDING AND BONDED STRUCTURE					
First Named Inventor/Applicant Name:	Qin-Yi Tong					
Filer:	Pau	ıl Steven Stellman/.	Jonathan Pan			
Attorney Docket Number:	TSS	SRA.012C1				
Filed as Large Entity						
Filing Fees for Utility under 35 USC 111(a)						
Description		Fee Code	Quantity	Amount	Sub-Total in USD(\$)	
Basic Filing:						
Pages:						
Claims:						
Miscellaneous-Filing:						
Petition:						
Patent-Appeals-and-Interference:						
Post-Allowance-and-Post-Issuance:						
Utility Appl Issue Fee		1501	1	960	960	

Description	Fee Code	Fee Code Quantity		Sub-Total in USD(\$)
Extension-of-Time:				
Miscellaneous:				
	Total in USD (\$)		960	

Electronic Acknowledgement Receipt				
EFS ID:	25885232			
Application Number:	14957501			
International Application Number:				
Confirmation Number:	1033			
Title of Invention:	METHOD FOR LOW TEMPERATURE BONDING AND BONDED STRUCTURE			
First Named Inventor/Applicant Name:	Qin-Yi Tong			
Customer Number:	135980			
Filer:	Paul Steven Stellman/Chelsea Burdeno			
Filer Authorized By:	Paul Steven Stellman			
Attorney Docket Number:	TSSRA.012C1			
Receipt Date:	25-MAY-2016			
Filing Date:	02-DEC-2015			
Time Stamp:	17:33:10			
Application Type:	Utility under 35 USC 111(a)			

Payment information:

Submitted with Payment	yes
Payment Type	Credit Card
Payment was successfully received in RAM	\$960
RAM confirmation Number	4321
Deposit Account	111410
Authorized User	KNOBBE MARTENS OLSON AND BEAR

The Director of the USPTO is hereby authorized to charge indicated fees and credit any overpayment as follows:

Charge any Additional Fees required under 37 CFR 1.16 (National application filing, search, and examination fees)

File Listing:					
Document Number	Document Description	File Name	File Size(Bytes)/ Message Digest	Multi Part /.zip	Pages (if appl.)
1	Issue Fee Payment (PTO-85B)	2016-05-25_IssueFeeTransmitt al_TSSRA-012C1.pdf	84957	no	1
'	issue ree rayment (170 055)		743507b930368d6840e71e29659b6380fe8 a23a8	110	
Warnings:					
Information:					
2	Miscellaneous Incoming Letter	2016-05-25_Comments_TSSRA -012C1.pdf	71533	no	2
	Miscellaneous meoning eccer		477fc556dfce772e5603a393e2bfc687e896 6ad6	110	_
Warnings:					
Information:					
3 Fee Worksheet (SB06)	Fee Worksheet (SB06)	fee-info.pdf	30804	no	2
	Tee Worksheet (5500)	2914d1c6c0d97c24513dc52d998a980af6a c8e16		_	
Warnings:					
Information:					
		Total Files Size (in bytes)	18	37294	

This Acknowledgement Receipt evidences receipt on the noted date by the USPTO of the indicated documents, characterized by the applicant, and including page counts, where applicable. It serves as evidence of receipt similar to a Post Card, as described in MPEP 503.

New Applications Under 35 U.S.C. 111

If a new application is being filed and the application includes the necessary components for a filing date (see 37 CFR 1.53(b)-(d) and MPEP 506), a Filing Receipt (37 CFR 1.54) will be issued in due course and the date shown on this Acknowledgement Receipt will establish the filing date of the application.

National Stage of an International Application under 35 U.S.C. 371

If a timely submission to enter the national stage of an international application is compliant with the conditions of 35 U.S.C. 371 and other applicable requirements a Form PCT/DO/EO/903 indicating acceptance of the application as a national stage submission under 35 U.S.C. 371 will be issued in addition to the Filing Receipt, in due course.

New International Application Filed with the USPTO as a Receiving Office

If a new international application is being filed and the international application includes the necessary components for an international filing date (see PCT Article 11 and MPEP 1810), a Notification of the International Application Number and of the International Filing Date (Form PCT/RO/105) will be issued in due course, subject to prescriptions concerning national security, and the date shown on this Acknowledgement Receipt will establish the international filing date of the application.



United States Patent and Trademark Office

UNITED STATES DEPARTMENT OF COMMERCE United States Patent and Trademark Office Address: COMMISSIONER FOR PATENTS P.O. Box 1450 Alexandria, Virginia 22313-1450 www.uspto.gov

APPLICATION	FILING or	GRP ART				
NUMBER	371(c) DATE	UNIT	FIL FEE REC'D	ATTY.DOCKET.NO	TOT CLAIMS	IND CLAIMS
14/957 501	12/02/2015	2895	3200	TSSRA 012C1	33	4

135980 Knobbe, Martens, Olson & Bear, LLP 2040 main Street Fourteenth Floor Irvine, CA 92614 CONFIRMATION NO. 1033 CORRECTED FILING RECEIPT



Date Mailed: 06/09/2016

Receipt is acknowledged of this non-provisional patent application. The application will be taken up for examination in due course. Applicant will be notified as to the results of the examination. Any correspondence concerning the application must include the following identification information: the U.S. APPLICATION NUMBER, FILING DATE, NAME OF APPLICANT, and TITLE OF INVENTION. Fees transmitted by check or draft are subject to collection. Please verify the accuracy of the data presented on this receipt. If an error is noted on this Filing Receipt, please submit a written request for a Filing Receipt Correction. Please provide a copy of this Filing Receipt with the changes noted thereon. If you received a "Notice to File Missing Parts" for this application, please submit any corrections to this Filing Receipt with your reply to the Notice. When the USPTO processes the reply to the Notice, the USPTO will generate another Filing Receipt incorporating the requested corrections

Inventor(s)

Qin-Yi Tong, Durham, NC, Deceased; Gaius Gillman Fountain JR., Youngsville, NC; Paul M. Enquist, Cary, NC;

Applicant(s)

ZIPTRONIX, INC., Morrisville, NC;

Power of Attorney: The patent practitioners associated with Customer Number <u>135980</u>

Domestic Priority data as claimed by applicant

This application is a CON of 14/754,111 06/29/2015 PAT 9331149

which is a DIV of 14/197,070 03/04/2014 PAT 9082627

which is a DIV of 13/341,273 12/30/2011 ABN

which is a CON of 12/954,740 11/26/2010 PAT 8153505

which is a CON of 12/720,368 03/09/2010 PAT 7871898

which is a CON of 11/980,664 10/31/2007 PAT 7807549

which is a CON of 10/913,441 08/09/2004 PAT 7387944

which is a CON of 09/505,283 02/16/2000 PAT 6902987

Foreign Applications for which priority is claimed (You may be eligible to benefit from the **Patent Prosecution Highway** program at the USPTO. Please see http://www.uspto.gov for more information.) - None. Foreign application information must be provided in an Application Data Sheet in order to constitute a claim to foreign priority. See 37 CFR 1.55 and 1.76.

Permission to Access Application via Priority Document Exchange: Yes

page 1 of 4

Permission to Access Search Results: Yes

Applicant may provide or rescind an authorization for access using Form PTO/SB/39 or Form PTO/SB/69 as appropriate.

If Required, Foreign Filing License Granted: 12/11/2015

The country code and number of your priority application, to be used for filing abroad under the Paris Convention,

is US 14/957,501

Projected Publication Date: Not Applicable

Non-Publication Request: No Early Publication Request: No

Title

METHOD FOR LOW TEMPERATURE BONDING AND BONDED STRUCTURE

Preliminary Class

257

Statement under 37 CFR 1.55 or 1.78 for AIA (First Inventor to File) Transition Applications: No

PROTECTING YOUR INVENTION OUTSIDE THE UNITED STATES

Since the rights granted by a U.S. patent extend only throughout the territory of the United States and have no effect in a foreign country, an inventor who wishes patent protection in another country must apply for a patent in a specific country or in regional patent offices. Applicants may wish to consider the filing of an international application under the Patent Cooperation Treaty (PCT). An international (PCT) application generally has the same effect as a regular national patent application in each PCT-member country. The PCT process **simplifies** the filing of patent applications on the same invention in member countries, but **does not result** in a grant of "an international patent" and does not eliminate the need of applicants to file additional documents and fees in countries where patent protection is desired.

Almost every country has its own patent law, and a person desiring a patent in a particular country must make an application for patent in that country in accordance with its particular laws. Since the laws of many countries differ in various respects from the patent law of the United States, applicants are advised to seek guidance from specific foreign countries to ensure that patent rights are not lost prematurely.

Applicants also are advised that in the case of inventions made in the United States, the Director of the USPTO must issue a license before applicants can apply for a patent in a foreign country. The filing of a U.S. patent application serves as a request for a foreign filing license. The application's filing receipt contains further information and guidance as to the status of applicant's license for foreign filing.

Applicants may wish to consult the USPTO booklet, "General Information Concerning Patents" (specifically, the section entitled "Treaties and Foreign Patents") for more information on timeframes and deadlines for filing foreign patent applications. The guide is available either by contacting the USPTO Contact Center at 800-786-9199, or it can be viewed on the USPTO website at http://www.uspto.gov/web/offices/pac/doc/general/index.html.

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countries such as China, Korea and Mexico. For questions regarding patent enforcement issues, applicants may call the U.S. Government hotline at 1-866-999-HALT (1-866-999-4258).

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ISSUE DATE ATTORNEY DOCKET NO. CONFIRMATION NO. APPLICATION NO. PATENT NO. 14/957,501 07/12/2016

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TSSRA.012C1

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06/22/2016

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ISSUE NOTIFICATION

The projected patent number and issue date are specified above.

Determination of Patent Term Adjustment under 35 U.S.C. 154 (b)

(application filed on or after May 29, 2000)

The Patent Term Adjustment is 0 day(s). Any patent to issue from the above-identified application will include an indication of the adjustment on the front page.

If a Continued Prosecution Application (CPA) was filed in the above-identified application, the filing date that determines Patent Term Adjustment is the filing date of the most recent CPA.

Applicant will be able to obtain more detailed information by accessing the Patent Application Information Retrieval (PAIR) WEB site (http://pair.uspto.gov).

Any questions regarding the Patent Term Extension or Adjustment determination should be directed to the Office of Patent Legal Administration at (571)-272-7702. Questions relating to issue and publication fee payments should be directed to the Application Assistance Unit (AAU) of the Office of Data Management (ODM) at (571)-272-4200.

APPLICANT(s) (Please see PAIR WEB site http://pair.uspto.gov for additional applicants):

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