

Evidence for Involvement of BRCA1 in Sporadic Breast Carcinomas

Craig S. Cropp, Heli A. Nevanlinna, Seppo Pyrhönen, Ulf-Håkan Stenman, Paul Salmikangas, Hans Albertsen, Ray White, and Robert Callahan

Laboratory of Tumor Immunology and Biology, National Cancer Institute, NIH, Bethesda, MD 20892 [C. C., R. C.]; Departments I and II of Obstetrics and Gynecology [H. A. N., U.-H. S., P. S.] and Radiotherapy and Oncology [S. P.], Helsinki University Central Hospital, Haartmaninkatu 2, 00290 Helsinki, Finland; and Eccles Institute of Human Genetics and Howard Hughes Medical Institute, University of Utah, Salt Lake City, Utah 84112 [H. A., R. W.]

Abstract

The hereditary breast cancer gene *BRCA1* previously has been localized to chromosome 17q21. We looked for evidence of involvement of this region of chromosome 17 in 130 sporadic breast cancers. Seventeen polymorphic sequence tagged site markers were examined in these tumors between the *D17S250* and *D17S579* loci to screen for deletions as measured by loss of heterozygosity. The smallest common region that was deleted occurred in the approximately 120-kilobase interval between the *D17S846* and *D17S746* loci within the *BRCA1* region. Delineation of this commonly deleted area should accelerate attempts to identify the involved gene(s) and its relationship to *BRCA1*.

Introduction

It is widely believed that there are one or more tumor suppressor genes on chromosome 17q for breast carcinoma (1-3). One candidate gene, *BRCA1*, is genetically linked to the development of some familial breast cancers and is located at 17q21 (4). Ovarian cancer is also known to have a hereditary component (5-7). Furthermore, a woman with ovarian cancer is at increased risk for developing breast cancer as a second primary tumor, and *vice versa* (8-10). Recently, the syndrome for familial breast/ovarian cancer has also been linked to the same region as *BRCA1* (11). The observed LOH¹ affecting the wild type chromosome in tumors from affected breast/ovarian cancer patients (12, 13) is consistent with the hypothesis that *BRCA1* is a tumor suppressor gene. However, whether the target genes for familial breast and breast/ovarian cancers are the same or are different closely linked genes is not known. In a variety of hereditary neoplasias the affected gene (e.g., *RB*, *TP53*, *APC*, etc.) has also been found to be frequently mutated in sporadic forms of the disease (14, 15). Previously, we (1) and others (2, 3, 16) have shown that sporadic breast carcinomas are frequently affected by LOH in the general region of *BRCA1*. The availability of additional polymorphic STS which have been genetically mapped to the region of chromosome 17q21 containing *BRCA1* (17) has provided us the opportunity to further define the region of this portion of the genome which is affected by LOH in sporadic primary human breast tumors. When this study was begun the consensus among published reports on the genetic and physical boundaries of *BRCA1* suggested that the centromeric boundary was *D17S250* (18-20). The telomeric boundary was less clear. In some reports it was *D17S588* while in others it was the more centromeric locus, *D17S579* (19, 21, 22). In our study we chose *D17S579* as our telomeric border. The interval between *D17S250* and *D17S579* is estimated to be roughly 3500 kilobases (23). In the present study we

describe a high density deletion map using 17 PCR-based polymorphic STS markers located between the *D17S250* and *D17S579* loci in 130 primary human breast tumors.

Materials and Methods

Primary breast carcinomas and matching peripheral lymphocytes were collected at the Helsinki University Central Hospital in Helsinki, Finland, from 130 patients who had received no prior therapy.

Genomic DNA was extracted and diluted to 100-200 ng/ μ l. PCR was performed with 100-200 ng template DNA, 10 mM Tris-HCl, 1.5 mM MgCl₂, 50 mM KCl, gelatin 0.1 mg/ml, 250 μ M concentration of each nucleoside triphosphate, 0.4 unit Taq polymerase (Boehringer Mannheim), and 30 pmol of each primer in a total volume of 25 μ l. The PCR product was identified by end labeling primers with [γ -³²P]ATP. All PCR reactions were performed on a GeneAmp PCR System 9600 starting with denaturation for 6 min at 94°C followed by 25 cycles of denaturation at 94°C for 10 s, annealing temperature for 10 s, and extension at 72°C for 20 s. The primers used, their annealing temperatures, and references are shown in Table 1.

The PCR products were diluted with loading buffer (95% formamide, 20 mM EDTA, 0.05% bromophenol blue, and 0.05% xylene cyanol), heat denatured and rapidly cooled. Samples were run in pairs (tumor and lymphocyte PCR product from the same patient) on a denaturing gel (7% acrylamide-32% formamide-6 M urea-1 $\times$ Tris-buffered EDTA) at a constant 65-70 W. After electrophoresis the gel was transferred to Whatman No. 3MM paper and autoradiography was performed with Kodak X-Omat AR film at -70°C.

Results and Discussion

We have previously defined three regions of chromosome 17q that are frequently affected by LOH in primary human breast tumors (1). In that study a putative target gene(s) was suggested in the interval between *D17S73* and *NME1* on chromosome 17q12-q21. In the present study we have examined an additional 17 polymorphic STS markers between *D17S250* and *D17S579* which is a subregion within the *D17S73* and *NME1* interval. This represents an average of one polymorphism every 210 kilobases since the distance between *D17S250* and *D17S579* is estimated to be 3500 kilobases. The total number of tumors examined, the percentage of the total number of tumors that were informative, and the percentage of the informative tumors which were deleted (i.e., LOH) is shown in Table 1. The overall frequency of LOH varies from 12 to 32% and is the highest between *D17S846* and *D17S776*.

To further define the region containing the putative target gene we have analyzed the genotypes of individual primary breast tumor DNAs for evidence of LOH. Shown in Fig. 1 are autoradiographs of the STS markers of four breast tumors between *D17S702* and *D17S856*. Tumor 20 showed no LOH at *D17S702* and *D17S746* but was deleted for the upper allele of *GAS* and the lower allele of *D17S846*. Markers *D17S776* and *D17S856* were not informative in this tumor. In tumor 26 no LOH was detected at *D17S702*, *D17S846*, and *D17S856*, but the tumor DNA was deleted for the lower allele of

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¹ The abbreviations used are: LOH, loss of heterozygosity; STS, sequence tag sites; PCR, polymerase chain reaction

Table 1 Frequency of LOH at chromosome 17q21 loci

The locus number of the STS marker, the PCR annealing temperature, and a reference is listed for each marker used. The total number of tumors examined for each marker is indicated by *N*. The percentage of the total number of tumors (*N*) that were informative is shown as %, and the numbers in parentheses are the expected heterozygosity for that STS marker. The percentage of the informative tumors (*I*) that were deleted is indicated by %LOH.

Locus/STS	Annealing temperature (°C)	Ref.	<i>N</i>	% I	% of LOH
<i>D17S250</i>	55	24	101	80 (81)	21
<i>THRA1</i>	55	25	95	39 (81)	27
<i>D17S700</i>	60	17	100	54 (44)	19
<i>D17S857</i>	55	26	102	67 (NP)	25
<i>D17S702</i>	52	17	102	90 (88)	23
<i>GAS</i>	60	27	97	52 (NP)	26
<i>D17S846</i>	55	28	95	78 (84)	32
<i>D17S746</i>	58	17	108	33 (44)	28
<i>D17S776</i>	58	29	105	60 (55)	30
<i>D17S856</i>	55	26	103	51 (NP)	19
<i>D17S648</i>	60	17	107	36 (29)	18
<i>D17S855</i>	55	26	60	50 (NP)	27
<i>D17S902</i>	60	17	50	74 (78)	27
<i>D17S859</i>	55	26	49	35 (NP)	12
<i>D17S750</i>	65	17	103	57 (66)	24
<i>D17S183</i>	55	30, 31	103	35 (40)	22
<i>D17S579</i>	55	32	109	86 (87)	13

D17S746. Tumor 63 was informative and unaltered at the *D17S702*, *GAS*, *D17S846*, and *D17S856* loci but was deleted for the upper allele of *D17S776*. In this patient the STS marker *D17S746* was not informative. Tumor 117 was informative and unaltered at the *D17S702*, *D17S776*, and *D17S856* loci but was deleted for the upper allele of *D17S846*. In this patient *D17S746* was not informative.

The genotypes of these four tumors and seven additional tumors for loci lying between *D17S250* and *D17S579* is shown in Fig. 2. Ten of the tumors examined in this study have clearly defined interstitial deletions within the *BRCA1* interval, while tumor 127 appears to have lost a large portion of the long arm of chromosome 17. In this study a total of nine tumors appeared to have lost an entire chromosome 17q arm, since all markers in the *BRCA1* region as well as telomeric markers, such as *D17S4*, were deleted. Six other tumors had LOH at all the markers in the *BRCA1* region but were informative and retained heterozygosity at *D17S4*. These data taken together indicate that the smallest commonly deleted region is located between *D17S846* and *D17S746*. This conclusion is both consistent with and extends previous studies in which fewer loci were examined within the *BRCA1* region (2, 3, 16). *D17S846* and *D17S746* are located on two overlapping recombinant P1 bacteriophage clones. The distance between them is estimated to be 120–150 kilobases based on physical mapping data² and could possibly contain several undiscovered genes.

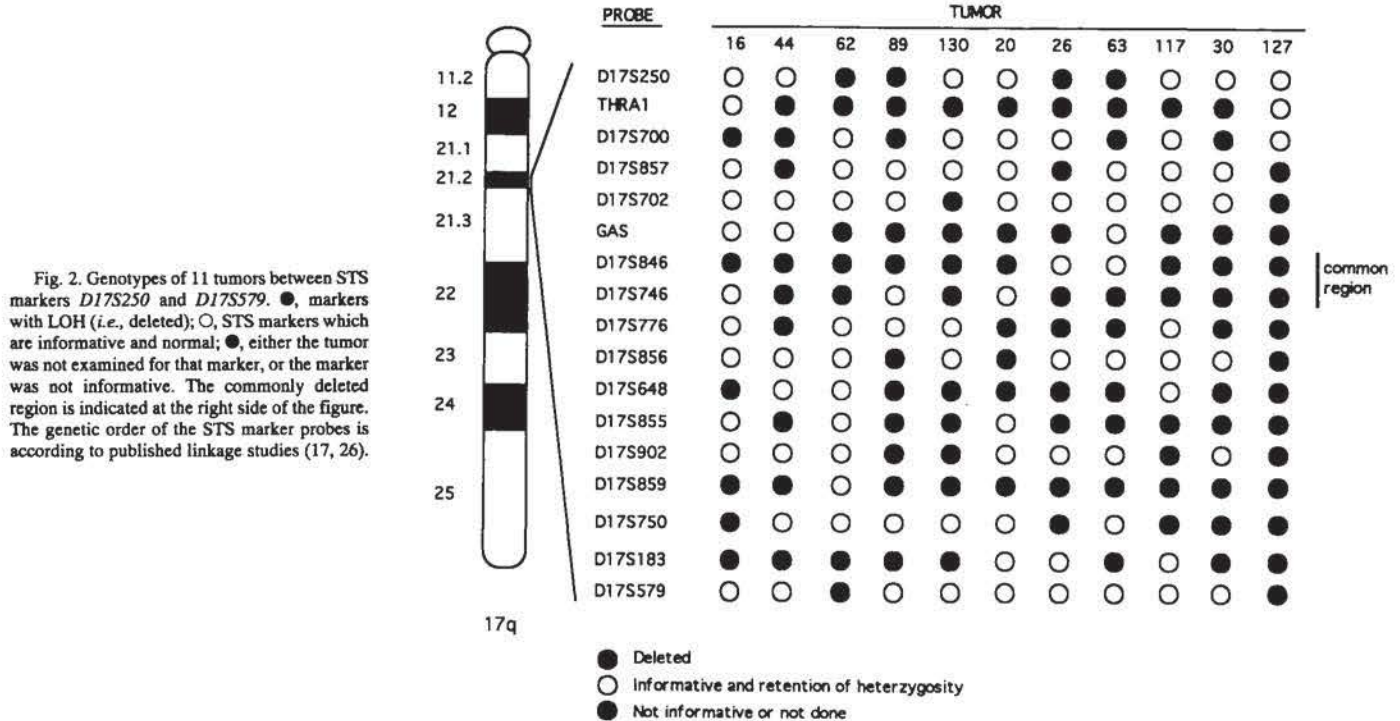
The increased availability of polymorphic STS markers in the region of *BRCA1* has also led to a continued shrinkage of the size of this region through genetic linkage studies in breast and breast/ovarian cancer families. Thus, in successive studies the telomeric border has moved centromerically from *D17S579* (18, 20, 21) to *D17S183* (located between *D17S750* and *D17S579*; Fig. 2) (26) to *D17S78* (located between *D17S750* and *D17S183*; Fig. 2) (33). Similarly the centromeric border has moved in the telomeric direction from *D17S250* (19, 20, 31) to *THRA1* (21, 30) to *D17S857* (13) to *D17S702* (34). The region we have defined as containing the target gene for LOH on chromosome 17q21 in sporadic breast tumors is compatible with these linkage studies defining *BRCA1* in familial breast and breast/ovarian cancer families. However, there is one report (29) which is potentially inconsistent with this conclusion. They show that in one family having the familial breast/ovarian cancer syndrome, one

affected member had a recombinational event telomeric of the locus *D17S776*. This locus is more telomeric than *D17S746* which is the telomeric boundary of the affected region defined in our study (Fig. 2). There are at least three explanations for this apparent paradox: (a) the locus we have defined may be relevant only in sporadic breast cancer and not in hereditary breast cancer; (b) possibly there are two closely linked genes which are independently altered in breast-only versus breast/ovarian families. A comparison of the linkage studies done in breast-only families versus those done in breast/ovarian families show that in the former group the altered gene lies between *THRA1* and *D17S579* (35), whereas in the latter group the location of the altered gene lies distal to *D17S776* and proximal to *D17S78* (29, 32). The region we have defined is consistent with the current linkage analysis of the location of the altered gene in the breast-only families; (c) our results are not incompatible with the linkage data of Goldgar *et al.* (29). However, this would require that the *BRCA1* gene is large (1–2 megabases) and that the recombinational event described by Goldgar *et al.* occurred within the gene. There are precedents for each of these requirements. For instance, the *DCC* gene on chromo-



Fig. 1. Autoradiographs of four tumors for the six STS polymorphic markers between *D17S702* and *D17S856*. Arrows, deleted allele indicating LOH. Tumor 20 is deleted for *GAS* and *D17S846*; tumor 26 is deleted for *D17S746*; tumor 63 is deleted for *D17S776*; and tumor 117 is deleted for *D17S846*. T, tumor; L, lymphocyte.

² H. Albertsen *et al.*, manuscript in preparation.



some 18q is approximately 1.4 megabases (36), and the Duchenne muscular dystrophy (*DMD*) gene on Xp21 is approximately 2.3 megabases (37). In the search for the *DMD* gene, linkage analysis was somewhat confusing because of the frequency of recombination events within such a large gene (38–40). Clearly, identification of the target gene(s) in sporadic breast carcinomas as well as the familial breast and breast/ovarian carcinoma loci will be required to distinguish between these possibilities.

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