NDC† 76439-205-90 (90 ct. bottle / 90 tablets)

## L-Methylfolate Calcium 15 mg **Tablets**

Ry

Prescription Dietary Supplement

DESCRIPTION: L-Methylfolate Calcium 15 mg is an orally DESCRIPTION: L-Methytolate Calcum 15 mg is an orally administered prescription dietary supplement specifically formulated for the dietary management of patients with unique nutritional needs requiring increased folate levels.

L-Methytfolate Calcium 15 mg should be administered under the supervision of a licensed medical practitioner.

Each coated, oblong, orange tablet contains the following dietary

Supplement Facts	Serving Size: 1 toblet	Servings For Container: 90
	Imount Per Serving	% Daily Value
L-Mathylfelate Caldium*, or 6(S)-S-MTHF-Co		

(and ct. bottle / 90 tablets)

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\*contains less than 1% d-methylfolate.
\*\*Daily Value not established for patients with unique nutritional needs who are in need of supplementation as directed by a licensed medical practitioner.

Other ingredients: Croscarmellose sodium, dicalcium phosphate, magnesium stearate, microcrystalline cellulose, silicon dioxide, stearia edi, and film coat (FD&C red #2, FD&C yellow #6 lake, polyethylene glycol, polyvinyl alcohol, talc, and titanium dioxide).

This product contains FD&C Yellow #6 Lake.

FOLATE REGULATION: The term "folate" are B vitamins that include FOLAITE REGULATION: That earn loater are of warming such at loader folic acid and any forms of active oteroylglutamates regardless of the reduction state of the molecule. Folates, or vitamin B9, are primarily hydrolyzed in the intestinal jejunum and the liver to the active circulating form of folate, i-methylfolate, with an intermediate stable form, 5,10-methylenetetrahydrofolate.

Individuals with genetic polymorphisms for the genes coding methylenetetrahydrofolate reductase (NTHFR) may not be capable of utilizing or metabolizing folic acid adequately for the vitamin B<sub>12</sub> dependent methylation cycle.

Folic acid, including reduced forms\* such as folinic acid, mo obscure pernicious anemia above 0.1 mg doses, and must administered under the supervision of a licensed medical

The 1971, 1972, 1973, 1980, 1984, 2000, and 2010 Federal Register Notices addressed this concern while establishing that increased folate was proper therapy in megalobiastic anemias - specifically where homocysteine levels were elevated or risk of neural tube defects (NTDs) was at issue. The Federal Register Notice of August 2, 1973 (38 FR 20750) specifically states that:

Dietary supplement preparations are available without a prescription (21 CFR 121.1134). Levels higher than dietary supplement amounts are available only with a prescription.

‡ It is not known whether or not I-methylfolate can obscure pernicious anemia above 0.1 mg doses, so caution is advised also with this form of folate.

Folic acid - including reduced forms, may be added to medical foods as defined in section 5(b)(3) of the Orphan Drug Act (21 USC 360ee(b)(3)), or to food (21 CFR 172.345).

INDICATIONS AND USAGE: L-Methylfolate Calcium 15 mg is indicated for the distinct nutritional requirements of patients in need of dietary supplementation as determined by a licensed medical practitioner. L-Methylfolate Calcium 15 mg should be administered under the supervision of a licensed medical practitioner.

CONTRAINDICATIONS: This product is contraindicated in patients with a known hypersensitivity to any of the ingredients.

WARNINGS: Caution is recommended in patients with a history of

PRECAUTIONS: General: Folate, when administered as a single agent in doses about 0.1 mg daily, may obscure the detection of vitamin B<sub>12</sub> deficiency (specifically, the administration of folic acid may reverse the hematological manifestations of B<sub>12</sub> deficiency, including pernicious anemia, while not addressing the neurological manifestations). Folate therapy alone is inadequate for treatment of a vitamin B<sub>12</sub> deficiency.

A major depressive episode may be the initial presentation of bipolar disorder. It is generally believed, (although not established in controlled trials) that treating such an episode with an antidepressant alone may increase the likelihood of a precipitation of a mixed/manic episode in patients at risk for bipolar disorder. L-Methylfolate Calcium 15 mg is not an antidepressant; however, folate has been showned enhance antidepressants effects of known antidepressants. Caution is recommended in patients with a history of bipolar illness. Patients with depressive symptoms should be adequately screened to determine if they are at risk for bipolar disorder since mood elevation in this population is possible.

PATIENT INFORMATION: L-Methylfolate Calcium 15 mg is a prescription dietary supplement to be used only under licensed medical supervision.

DRUG INTERACTIONS: Drugs which may interact with foliate include:
 Antiepileptic drugs (AED): The AED class including, but not limited to, phenyloin, carbamazepine, primidone, valproic acid, fosphenyloin, valproate, phenobarbital and lamotrigine have been shown to impair foliate absorption and increase the metabolism of circulating foliate.

Additionally, concurrent use of folia acid has been associated.

 Additionally, concurrent use of folic acid has been as with enhanced phenytoin metabolism, lowering the level of the AED in the blood and allowing breakthrough seizures to occur. Caution should be used when prescribing this product among patients who are receiving treatment with phenytoin and other autions wileste.

Capecitabine: Folinic acid (5-formyltetrahydrofolate) may

increase the toxicity of Capecitabine.

Cholestyramine: Reduces folic acid absorption and reduces

Cholestyramline: Réduces folic acid absorption and reduces serum folate levels.
 Colestipol: Reduces folic acid absorption and reduces serum folate levels.
 Colestipol: Reduces folic acid absorption and reduces serum folate levels.
 Dilydrofolate Reduces folic acid absorption and reduces serum folate levels.
 Dilydrofolate Reductase Inhibitors (DHFRI): DHFRIs block the conversion of folic acid to its active forms, and lower plasma and red blood cell folate levels. DHFRIs include aminopterin, methotrexate, pyrimethamine, triamterene, and trimethoprim.
 Fluoxetime: Fluoxetime exerts a noncompetitive inhibition of the 5-methyltetrahydrofolate active transport in the intestine.
 Isotretinion: Reduced folate levels have occurred in some patients taking isotretinion.

patients taking isotretinoin.
 L-dopa, triamterene, colchicine, and trimethoprim may decrease

plasma folate levels.

Nonsteroidal Anti-inflammatory Drugs (NSAIDs): NSAIDs have

Nonsteroidal Anti-inflammatory Drugs (NSAIDs): NSAIDs have been shown to inhibit some foldet dependent enzymes in laboratory experiments. NSAIDs include buporfern, approxen, indomethacin and sulindac.

Oral Contraceptives therapy.

Methylprednisolone: Reduced serum folate levels have been noted after treatment with methylprednisolone.

Pancreatic Enzymes: Reduced folate levels have occurred in some patients taking pancreatic extracts, such as pancreatin and pancrelipase.

Pentamidine: Reduced folate levels have been seen with prolonged intravenous pentamidine.

Pyrimethamine: High levels of folic acid may result in decreased serum levels of pyrimethamine.

Smoking and Alcohof: Reduced serum folate levels have been noted.

Sulfasalazine: Inhibits the absorption and metabolism of folic acid.

Metformin treatment in patients with type 2 diabetes decreases serum folate.

serum folate. Warfarin can produce significant impairment in folate status after

Warfarn can produce significant impairment in rolate status after a 6-month hierapy.
 Folinic acid may enhance the toxicity of Tworouracil.
 Concurrent administration of chloramphenicol and folinic acid in folate-deficient patients may result in antalgonism of the haematopoietic response to folate.
 Caution should be exercised with the concomitant use of folinic acid and trimethoprim-sulfamethoxazole for the acute treatment of Pneumocystic acrinii pneumonia in patients with HIV infection as it is associated with increased rates of treatment fallure and mortality in a placebo controlled study.

PREGNANCY and NURSING MOTHERS: L-Methylfolate Calcium 15 mg is not intended for use as a prenatal/postnatal multivitamin for lactating and non-lactating mothers. This product contains a B vitamin in reduced form. Talk with your medical practitioner before using if pregnant or lactating.

ADVERSE REACTIONS: Allergic sensitization has been reported following both oral and parental administration of folic acid, and may possibly occur with other forms of folate.

DOSAGE AND ADMINISTRATION: The usual adult dose is 15 mg daily with or without food or as directed by a licensed medical practitioner.

HOW SUPPLIED: L-Methylfolate Calcium 15 mg tablets are coated, oblong, orange tablets debossed "BP" on top and "1000" on bottom, and are supplied in bottles of 90 tablets.

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1 These products are dietary supplements that – due to increased folate levels (8/2/73 38 FR 20750), require an Rx on the label because of increased risk associated with masking of 8/12 deficiency. As such, this product requires licensed medical supervision, an Rx status, and a National Drug Code (NDC) as required by pedigree reportion requirements.

STORAGE: Store at controlled room temperature 15°-30°C (59°-86°F). (See USP). Protect from light and moisture. Dispense in a tight, light-resistant container.

KEEP THIS OUT OF REACH OF CHILDREN.
All prescriptions using this product shall be pursuant to state statutes as applicable. This is not an Orange Book product.
Call your medical practitioner about side effects,
You may report side effects by calling 813-283-1344.

L-Methylfolate Calcium 15 mg Tablets

Prescription Dietary Supplement 90 ct. Bottle (90 tablets)

Manufactured for: Virtus Pharmaceuticals, LLC Tampa, FL 33619 www.virtusRX.com MADE IN USA



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