Prenate Restore

Rx postnatal vitamin with probiotics, DHA, and chelated iron

Rx Only Dietary Supplement

DESCRIPTION: PRENATE® Restore is a prescription postnatal vitamin dietary supplement that contains probiotics, DHA, and advanced calcium. Each dark blue softgel is imprinted with "Omega Rx" on one side and blank on the other.

Supplement Facts Serving Size: 1 Softgel Amount per Serving:		% Daily Value	% Daily Value (Pregnant & Lactating Women)
Vitamin C (as ascorbic acid)	85 mg	140%	140%
Vitamin D3 (as cholecalciferol)	25 mcg	130%	130%
Vitamin E (as dl-alpha tocopheryl acetate)	4.5 mg	30%	30%
Folate (as 1.11 mg of (6S)-N5-methyltetrahydrofolic acid calcium salt (equivalent to 600 mcg of folic acid) and folic acid, USP 400 mcg)	1700 mcg DFE	430%	280%
Vitamin B6 (as pyridoxine HCl)	25 mg	1250%	1000%
Vitamin B12 (as cyanocobalamin)	12 mcg	200%	150%
Biotin	500 mcg	170%	170%
Calcium (as a blend of Formical® (calcium formate) and calcium carbonate)	156 mg	20%	10%
Iron (as a blend of Sumalate® (ferrous asparto glycinate and ferrous fumarate)	e) 27 mg	150%	150%
Magnesium (as magnesium oxide)	45 mg	10%	10%
Bacillus Coagulans (as Lactospore®) 150 Million CFU	10 mg	†	†
Docosahexaenoic Acid (DHA)	400 mg	t	†
† Daily Value (DV) not established.			

OTHER INGREDIENTS: Beeswax, palm shortening, and soy lecithin. Gelatin capsule (FD&C Blue #1, FD&C Red #3, gelatin, glycerin, purified water, sorbitol, and titanium dioxide).

INDICATIONS: PRENATE® Restore is a multivitamin/multimineral fatty acid dietary supplement indicated for use in improving the nutritional status of women throughout pregnancy and in the postnatal period for both lactating and nonlactating mothers. PRENATE® Restore can also be beneficial in improving the nutritional status of women prior to conception.

CONTRAINDICATIONS: PRENATE® Restore is contraindicated in patients with a known hypersensitivity to any of the ingredients.

WARNING: Ingestion of more than 3 grams of omega-3 fatty acids (such as DHA) per day has been shown to have potential antithrombotic effects, including an increased bleeding time and International Normalized Ratio (INR), Administration of omega-3 fatty acids should be avoided in patients taking anticoagulants and in those known to have an inherited or acquired predisposition to bleeding.

PRECAUTIONS: Folic acid alone is improper therapy in the treatment of pernicious anemia and other megaloblastic anemias where Vitamin B12 is deficient. Folic acid in doses above 1.0 mg daily may obscure pernicious anemia in that hematologic remission can occur while neurological manifestations progress.

WARNING: Accidental overdose of iron-containing products is a leading cause of fatal poisoning in children under 6. Keep this product out of reach of children. In case of accidental overdose, call a doctor or poison control center immediately.

ADVERSE REACTIONS: Allergic sensitization has been reported following both oral and parenteral administration of folic acid.

DOSAGE AND ADMINISTRATION: One softgel daily, or as directed by a physician.

HOW SUPPLIED: Bottles of 30 softgels. (75854-308-30). The listed product number is not a National Drug Code. Instead, Avion Pharmaceuticals has assigned a product code formatted according to standard industry practice to meet the formatting requirements of pharmacy and healthcare insurance computer systems.

STORAGE: Store at 20° - 25°C (68° - 77°F); excursions permitted to 15° - 30°C (59° - 86°F).

[See USP Controlled Room Temperature.]

KEEP THIS AND ALL MEDICATIONS OUT OF THE REACH OF CHILDREN.

THESE STATEMENTS HAVE NOT BEEN EVALUATED BY THE FOOD AND DRUG ADMINISTRATION. THIS PRODUCT IS NOT Intended to diagnose, treat, cure, or prevent any disease.

MANUFACTURED FOR:

Avion Pharmaceuticals, LLC Alpharetta, GA 30005 1-888-61-AVION L-0170 Rev 0620-01

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Sumalate® is a registered trademark of Albion Laboratories, Inc., covered by one or more claims of U.S. Patent Nos. 5,516,925, 6,716,814, 8,007,846 and 8,425,956.

LactoSpore® is a registered trademark of Sabinsa Corporation.

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