

DESCRIPTION: PRENATE DHA® is a prescription prenatal/postnatal multivitamin/mineral/essential fatty acid softgel. Each softgel is blue in color, opaque, and imprinted with "DHA" on one side and blank on the other.

Amount per Serving:		% Daily Value	% Daily Value for Pregnant & Lactating Womer
Vitamin C (as ascorbic acid)	90 mg	150%	150%
Vitamin D3 (as cholecalciferol)	10 mcg	50%	70%
Vitamin E (as dl-alpha tocopherol acetate)	18 mg	120%	95%
Vitamin B6 (as pyridoxine HCI)	26 mg	1300%	1040%
Folate 170 (as (6S)-N5-methyltetrahydrofolic acid calciu (equivalent to 600 mcg of folic acid) and folic			280%
Vitamin B12 (as cyanocobalamin)	25 mcg	420%	310%
Calcium (as Formical® (calcium formate))	155 mg	15%	10%
Iron (as Sumalate® (ferrous asparto glycinate)) 18 mg	100%	100%
Magnesium (as magnesium oxide)	50 mg	15%	10%
Docosahexaenoic Acid (DHA)	300 mg	†	†
Decedent Autoriolo Fiora (DTIFT)	ooo mg		'

PRENATE DHA® contains fish oil and soy.

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

INDICATIONS: PRENATE DHA® 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 DHA® can also be beneficial in improving the nutritional status of women orior to conception.

CONTRAINDICATIONS: PRENATE DHA® 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: Before, during and/or after pregnancy, one softgel daily or as directed by a physician.

booket And Administration. Delote, during and/or after pregnancy, one softger daily of as directed by a physician

HOW SUPPLIED: Bottles of 30 softgels. (75854-312-30).

The listed product number is not a National Drug Code, but has instead merely been formatted to comply with standard industry practice for pharmacy and insurance computer systems.

Store at 20° - 25°C (68° - 77°F); excursions permitted to 15° - 30°C (59° - 86°F). [See USP Controlled Room Temperature.]

MANUFACTURED FOR:

Avion Pharmaceuticals, LLC Atlanta, GA 30005 1-888-61-AVION

L-0177 Rev 0620-02

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.

Formical® is a registered trademark of Nephro-Tech 1, LLC, covered by one or more claims of U.S. Patent No. 6,528,542. Sumalate® is a registered trademark of Albino 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