

30 CAPSULES

NDC 18860-752-01

natelle®ONE

Rx Prenatal Vitamin & Plant-Based DHA.
Now with 28 mg Iron

Rx Only

Dist by: **AZUR PHARMA INC.**
1818 Market Street, Suite 2350
Philadelphia, PA 19103
www.azurpharma.com

Mfg by: **Accucaps Industries Ltd**
Windsor, Ontario, Canada

Natelle® is a registered trademark of Pharmelle



Description: Natelle® One capsules for oral administration are supplied as oblong red soft gelatin capsules, imprinted with "Natelle®" in white ink.

Each capsule contains:

Docosahexaenoic Acid (DHA)	250 mg
Eicosapentaenoic Acid (EPA)	Not more than 0.625 mg
Calcium (Tricalcium Phosphate)	102 mg
Iron (Ferrous Fumarate)	28 mg
Vitamin C (Ascorbic Acid)	30 mg
Vitamin B-6 (Pyridoxine HCl)	25 mg
Vitamin E (D-Alpha Tocopherol)	30 IU
Folic Acid	1 mg

DHA is an omega-3 fatty acid. The DHA in Natelle® One is derived from the oils of microalgae.

Other Ingredients: Gelatin (Bovine), Beeswax, Glycerine, Soybean Oil, L edifin, Titanium Dioxide, Ethyl Vanillin.

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

Indications and Usage: Natelle® One is indicated to provide vitamin/mineral and omega-3 fatty acid supplementation to women throughout pregnancy, during the postnatal period for both lactating and non-lactating mothers, and throughout the childbearing years. Natelle® One may be beneficial in improving the nutritional status of women prior to conception.

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

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

Warnings: Ingestion of more than 3 grams of omega-3 fatty acids per day has been shown to have potential antithrombotic effects, including increased bleeding time and INR. Administration of omega-3 fatty acids should be avoided in patients on anticoagulants and in those known to have an inherited or acquired bleeding diathesis.

Precautions: Folic acid in doses above 0.1 mg daily may obscure pernicious anemia in that hematologic remission can occur while neurological manifestations remain progressive.

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

Dosage and Administration: One capsule daily or as directed by a physician.

Store at controlled room temperature 15°-30°C (59°-86°F).

To report a serious adverse event or obtain product information, contact (800) 890-3058.

NAT115-10-02



Lot/Exp.