PREGENNA- beta carotene, as corbic acid, cholecalciferol, .alpha.-tocopherol acetate, pyridoxine hydrochloride, biotin, folic acid, levomefolate calcium, cyanocobalamin, calcium carbonate, magnesium oxide, ferrous bisglycinate, and potassium iodide tablet Redmont Pharmaceuticals, LLC

PreGenna

71741-072-30

Prenatal/Postnatal Dietary Supplement

Dispensed by Prescription

DESCRIPTION

PreGenna is a prescription prenatal/postnatal multivitamin/multimineral dietary supplement.

Supplement Facts			
	% Daily Value		
300 mcg RAE	33%		
60 mg	67%		
10 mcg	50%		
4.5 mg (10 IU)	33%		
26 mg	1529%		
0.280 mg	933%		
1.67 mg DFE	418%		
0.67 mg DFE			
1 mg DFE			
0.013 mg	542%		
80 mg	6%		
25 mg	6%		
20 mg	111%		
0.150 mg	100%		
	300 mcg RAE 60 mg 10 mcg 4.5 mg (10 IU) 26 mg 0.280 mg 1.67 mg DFE 0.67 mg DFE 1 mg DFE 0.013 mg 80 mg 25 mg 20 mg		

^{**} Daily Value not established.

OTHER INGREDIENTS: Microcrystalline Cellulose, Maltodextrin, Croscarmellose Sodium, Silicon Dioxide, Stearic Acid, Magnesium Stearate, Film Coating (Hydroxypropyl Methylcellulose, Polyethylene Glycol, Titanium Dioxide, FD&C Blue # 1)

Allergen: NONE

INDICATIONS

PreGenna is a multivitamin/multimineral dietary supplement indicated for the use in improving the

nutritional status of women throughout pregnancy and in the postnatal period for both lactation on nonlactating mothers.

PreGenna can also be beneficial in improving the nutritional status of women prior to conception.

CONTRAINDICATIONS

PreGenna is contraindicated in patients with a known hypersensitivity to any of the ingredients.

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 poising in children under 6. Keep this product out of the reach of children. In the 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 tablet daily or as directed by a physician.

HOW SUPPLIED

Bottles of 30 tablets (71741-072-30¹). Tablet is light blue, oblong.

This product is a prescription-folate with or without other dietary ingredients that – due to increased folate levels increased risk associated with masking B12 deficiency (pernicious anemia) requires administration under the care of a licensed medical practitioner (64 FR 8760). 1-3 The most appropriate way to ensure pedigree reporting consistent with these regulatory guidelines and safety monitoring is to dispense this product only by prescription. This is not an Orange Book product. This product may be administered only under a physician's supervision and all prescriptions using this product shall be pursuant to state statues as applicable. The ingredients, indication or claims of this product are not to be construed to be drug claims.

- 1. Federal Register Notice of August 2, 1973 (38 FR 20750)
- 2. Federal Register Notice of October 17, 1980 (45 FR 69043, 69044)
- 3. Federal Register Notice of March 5, 1996 (61 FR 8760)

STORAGE AND HANDLING

STORAGE

Store at 20°-25° C (68°-77°F) excursions permitted to 15°-30°C (59°-86°F) [See USP Controlled Room Temperature.] Avoid excessive heat, light and moisture.

¹ Redmont Pharmaceuticals does not represent this product code to be National Drug Code (NDC). Product codes are formatted according to standard industry practice, to meet the formatting requirement by pedigree reporting and supply-chain control including pharmacies.

TAMPER EVIDENT

Do not use if seal is broken or missing.

MADE IN USA

Distributed by: Redmont Pharmaceuticals, LLC Birmingham, AL 35209 800-986-5909

Ferrochel™ is a trademark of Albion Laboratories, Inc.

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.

KEEP THIS AND ALL MEDICATIONS OF THE REACH OF CHILDREN.

PRINCIPAL DISPLAY PANEL - 30 Tablet Bottle Label

71741-072-30

PreGenna

PRENATAL/POSTNATAL
Dietary Supplement
30 TABLETS

Dispensed by Prescription



DOSAGE AND ADMINISTRATION:

Before, during and/or after pregnancy, one tablet daily or as directed by a physician.

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 the case of accidental overdose, call a doctor or poison control center immediately.

STORAGE: Store at 20°-25°C (68°-77°F) excursions permitted to 15°-30°C (59°-86°F) [See USP Controlled Room Temperature.] Avoid excessive heat, light and moisture.

71741-072-30

PreGenna

PRENATAL/POSTNATAL

Dietary Supplement

30 TABLETS

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Dispensed by Prescription

See package outsert for full prescribing information.



Distributed by: Redmont Pharmaceuticals, LLC Birmingham, AL 35209 (800) 986-5909 MADE IN USA

PREGENNA

beta carotene, ascorbic acid, cholecalciferol, .alpha.-tocopherol acetate, pyridoxine hydrochloride, biotin, folic acid, levomefolate calcium, cyanocobalamin, calcium carbonate, magnesium oxide, ferrous bisglycinate, and potassium iodide tablet

Product Information

Product Type DIETARY SUPPLEMENT Item Code (Source) NHRIC:71741-072

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
Beta Carotene (UNII: 01YAE03M7J) (Beta Carotene - UNII:01YAE03M7J)	Beta Carotene	300 ug	
Ascorbic Acid (UNII: PQ6CK8PD0R) (Ascorbic Acid - UNII:PQ6CK8PD0R)	Ascorbic Acid	60 mg	
Cholecalciferol (UNII: 1C6 V77QF41) (Cholecalciferol - UNII:1C6 V77QF41)	Cholecalciferol	10 ug	
.ALPHATO COPHEROL ACETATE (UNII: 9E8X80D2L0) (.ALPHATOCOPHEROL - UNII:H4N855PNZ1)	.ALPHATOCOPHEROL ACETATE	4.5 mg	
PYRIDO XINE HYDRO CHLO RIDE (UNII: 68 Y4CF58 BV) (PYRIDO XINE - UNII: KV2JZ1BI6Z)	PYRIDO XINE HYDRO CHLO RIDE	26 mg	
Biotin (UNII: 6SO6U10H04) (Biotin - UNII:6SO6U10H04)	Biotin	0.280 mg	
Folic Acid (UNII: 935E97BOY8) (Folic Acid - UNII:935E97BOY8)	Folic Acid	0.400 mg	
LEVOMEFOLATE CALCIUM (UNII: A9R10K3F2F) (LEVOMEFOLIC ACID - UNII:8S95DH25XC)	LEVOMEFOLATE CALCIUM	0.600 mg	
Cyanocobalamin (UNII: P6 YC3EG204) (Cyanocobalamin - UNII:P6 YC3EG204)	Cyanocobalamin	0.013 mg	
Calcium Carbonate (UNII: H0G9379FGK) (Calcium Cation - UNII:2M83C4R6ZB)	Calcium Carbonate	80 mg	
Magnesium Oxide (UNII: 3A3U0GI71G) (Magnesium Cation - UNII:T6V3LHY838)	Magnesium Oxide	25 mg	
FERROUS BISGLYCINATE (UNII: SFW1D987QV) (Ferrous Cation - UNII:GW89581OWR)	Ferrous Cation	20 mg	
Potassium Iodide (UNII: 1C4QK22F9J) (Iodide Ion - UNII:09G4I6V86Q)	Potassium Iodide	0.150 mg	

Inactive Ingredients	
Ingredient Name	Strength
MICRO CRYSTALLINE CELLULO SE 102 (UNII: PNR0 YF6 9 3 Y)	
Maltodextrin (UNII: 7CVR7L4A2D)	
Croscarmellose Sodium (UNII: M28 OL1HH48)	
Silicon Dioxide (UNII: ETJ7Z6XBU4)	
Stearic Acid (UNII: 4ELV7Z65AP)	
MAGNESIUM PALMITOSTEARATE (UNII: R40XA9G5BV)	
Hypromellose, Unspecified (UNII: 3NXW29V3WO)	
Polyethylene Glycol, Unspecified (UNII: 3WJQ0SDW1A)	
Titanium Dioxide (UNII: 15FIX9V2JP)	
FD&C Blue No. 1 (UNII: H3R47K3TBD)	

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NHRIC:71741-072-30	30 in 1 BOTTLE		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
DIETARY SUPPLEMENT		09/12/2019	

Serving Size:		Serving per Container :
	Amount Per Serving	% Daily Value
color		
scoring	1	
shape		
size (solid drugs)	19 mm	

Labeler - Redmont Pharmaceuticals, LLC (080843607)

Revised: 9/2019 Redmont Pharmaceuticals, LLC