

PRENATAL PLUS- vitamin a, ascorbic acid, cholecalciferol, .alpha.-tocopherol acetate, dl-, thiamine mononitrate, riboflavin, niacinamide, pyridoxine hydrochloride, folic acid, cyanocobalamin, calcium carbonate, iron, zinc oxide, and cupric oxide tablet
Nationwide Laboratories

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Prenatal Plus
Iron Tablets

Each Tablet Contains :	Amount
VITAMINS:	per Tablet
Vitamin A (from Acetate and Beta Carotene)	4000 IU
Vitamin C (Ascorbic Acid)	120 mg
Vitamin D-3 (Cholecalciferol)	400 IU
Vitamin E (dl-Alpha Tocopheryl Acetate)	22 IU
Thiamine (Vitamin B-1 from Thiamine Mononitrate)	1.84 mg
Riboflavin (Vitamin B-2)	3 mg
Niacin (as Niacinamide)	20 mg
Vitamin B-6 (from Pyridoxine HCl)	10 mg
Folic Acid	1 mg
Vitamin B-12 (Cyanocobalamin)	12 mcg
MINERALS:	
Calcium (from Calcium Carbonate)	200 mg
Iron (from Carbonyl Iron)	29 mg
Zinc (from Zinc Oxide)	25 mg
Copper (from Cupric Oxide)	2 mg

Other Ingredients

Ascorbyl Palmitate, Citric Acid anhydrous, DL- alpha Tocopherol, Ethylcellulose, FD&C Blue #2 lake, FD&C Red #40 lake, FD&C Yellow #5 lake, FD&C Yellow #6 lake, Glucose, Gum Acacia, Hypromellose, Magnesium Stearate, Maize Starch, Maltodextrin, Methylcellulose, Microcrystalline Cellulose, Mineral Oil, Mono- and di-glycerides, Polyethylene Glycol, Pregelatinized Corn Starch, Silicon Dioxide, Sorbic Acid, Soy Protein, Stearic Acid, Sucrose, Titanium Dioxide, Tricalcium Phosphate.

Indication

To provide Vitamin and Mineral supplementation throughout pregnancy and during the postnatal period for both the lactating and non-lactating mother. It is also useful for improving nutritional status prior to conception.

Dosage

As a dietary adjunct before, during and after pregnancy, take one tablet daily with a meal, or as directed by physician.

WARNING

Accidental overdose of iron-containing products is a leading cause of fatal poisoning in children under age 6. Keep this product out of the reach of children. In case of accidental overdose, call a doctor or Poison Control Center immediately.

Precautions

Folic acid may partially correct the hematological damage due to Vitamin B12 deficiency of pernicious anemia, while the associated neurological damage progresses.

This product contains FD&C Yellow No. 5 (tartrazine) which may cause allergic-type reactions (including bronchial asthma) in certain susceptible persons. Although the overall incidence of FD&C Yellow No. 5 (tartrazine) sensitivity in the general population is low, it is frequently seen in patients who also have aspirin hypersensitivity.

Store at room temperature. Dispense in a well closed light-resistant container, with a child resistant cap.

CALL YOUR DOCTOR FOR MEDICAL ADVICE ABOUT SIDE EFFECTS. YOU MAY REPORT SIDE EFFECTS TO THE FDA AT 1-800-FDA-1088 (TOLL FREE).

Keep this and all medication out of the reach of children.

Do not use if imprinted Safety Seal under cap is broken or missing.

Distributed by: **Nationwide Laboratories, Iselin NJ 08830**

Lot No.: Exp. Date: Rev. 7/12

PRINCIPAL DISPLAY PANEL - 100 Tablet Bottle Label

Nationwide Laboratories LLC

NDC 42937-705-10

**Prenatal Plus
Iron Tablets**

**Multivitamin/Multimineral
Supplement**

**For Use Before, During
and After Pregnancy.**

Rx only

100 Tablets

Prenatal Plus Iron Tablets

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100 Tablets

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Niacin (as Niacinamide)	20 mg
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Other Ingredients: Acetyl Palmitate, Citric Acid anhydrous, DL-alpha Tocopherol, Ethylcellulose, FD&C Blue #2 Lake, FD&C Red #40 Lake, FD&C Yellow #5 Lake, FD&C Yellow #6 Lake, Glucose, Gum Acacia, Hydroxypropyl Cellulose, Magnesium Stearate, Maltose Syrup, Maltodextrin, Methylcellulose, Microcrystalline Cellulose, Mineral Oil, Mono- and di-glycerides, Polyethylene Glycol, Pregelatinized Corn Starch, Silicon Dioxide, Sulfuric Acid, Soy Protein, Stearic Acid, Sucrose, Tannin Extract, Tricalcium Phosphate.

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Distributed by: Nationwide Laboratories, Inc. NJ 08830

Lot No.: Exp. Date: Res. 7/12



PRENATAL PLUS

vitamin a, ascorbic acid, cholecalciferol, .alpha.-tocopherol acetate, dl-, thiamine mononitrate, riboflavin, niacinamide, pyridoxine hydrochloride, folic acid, cyanocobalamin, calcium carbonate, iron, zinc oxide, and cupric oxide tablet

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:42937-705
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Vitamin A (UNII: 81G40H8B0T) (Vitamin A - UNII:81G40H8B0T)	Vitamin A	4000 [iU]
Ascorbic Acid (UNII: PQ6CK8PD0R) (Ascorbic Acid - UNII:PQ6CK8PD0R)	Ascorbic Acid	120 mg
Cholecalciferol (UNII: 1C6V77QF41) (Cholecalciferol - UNII:1C6V77QF41)	Cholecalciferol	400 [iU]
.Alpha.-Tocopherol Acetate, DL- (UNII: WRIWPIEW8) (.Alpha.-Tocopherol, DL- - UNII:7QWA1RIO01)	.Alpha.-Tocopherol, DL-	22 [iU]
Thiamine Mononitrate (UNII: 8K0I04919X) (Thiamine Ion - UNII:4ABT0J945J)	Thiamine	1.84 mg
Riboflavin (UNII: TLM2976OFR) (Riboflavin - UNII:TLM2976OFR)	Riboflavin	3 mg
Niacinamide (UNII: 25X51I8RD4) (Niacinamide - UNII:25X51I8RD4)	Niacinamide	20 mg
Pyridoxine Hydrochloride (UNII: 68Y4CF58BV) (Pyridoxine - UNII:KV2JZ1BI6Z)	Pyridoxine Hydrochloride	10 mg
Folic Acid (UNII: 935E97BOY8) (Folic Acid - UNII:935E97BOY8)	Folic Acid	1 mg
Cyanocobalamin (UNII: P6YC3EG204) (Cyanocobalamin - UNII:P6YC3EG204)	Cyanocobalamin	12 ug
Calcium Carbonate (UNII: H0G9379FGK) (Calcium Cation - UNII:2M83C4R6ZB, Carbonate Ion - UNII:7UJQ5OPE7D)	Calcium Carbonate	200 mg
Iron (UNII: E1UOL152H7) (Iron - UNII:E1UOL152H7)	Iron	29 mg
Zinc Oxide (UNII: SOI2LOH54Z) (Zinc Oxide - UNII:SOI2LOH54Z)	Zinc Oxide	25 mg
Cupric Oxide (UNII: V1XJQ704R4) (Cupric Cation - UNII:8CBV67279L)	Cupric Cation	2 mg

Inactive Ingredients

Ingredient Name	Strength
Ascorbyl Palmitate (UNII: QN83US2B0N)	

Anhydrous Citric Acid (UNII: XF417D3PSL)
.Alpha.-Tocopherol, DI- (UNII: 7QWA1RIO01)
ETHYLCELLULOSE, UNSPECIFIED (UNII: 7Z8S9VYZ4B)
FD&C Blue No. 2 (UNII: L06K8R7DQK)
Aluminum Oxide (UNII: LMI26O6933)
FD&C Red No. 40 (UNII: WZB9127XOA)
FD&C Yellow No. 5 (UNII: I753WB2F1M)
FD&C Yellow No. 6 (UNII: H77VEI93A8)
DEXTROSE, UNSPECIFIED FORM (UNII: IY9XDZ35W2)
Acacia (UNII: 5C5403N26O)
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)
Magnesium Stearate (UNII: 70097M6I30)
Starch, Corn (UNII: O8232NY3SJ)
Maltodextrin (UNII: 7CVR7L4A2D)
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)
Mineral Oil (UNII: T5L8T28FGP)
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)
Silicon Dioxide (UNII: ETJ7Z6XBU4)
Sorbic Acid (UNII: X045WJ989B)
Soy Protein (UNII: R44IWB3RN5)
Stearic Acid (UNII: 4ELV7Z65AP)
Sucrose (UNII: C151H8M554)
Titanium Dioxide (UNII: 15FIX9V2JP)
Tricalcium Phosphate (UNII: K4C08XP666)

Product Characteristics

Color	YELLOW (Tan)	Score	no score
Shape	OVAL	Size	18mm
Flavor		Imprint Code	CIS;28
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:42937-705-10	100 in 1 BOTTLE; Type 0: Not a Combination Product	06/01/2011	
2	NDC:42937-705-16	250 in 1 BOTTLE; Type 0: Not a Combination Product	06/01/2011	
3	NDC:42937-705-18	500 in 1 BOTTLE; Type 0: Not a Combination Product	06/01/2011	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
UNAPPROVED DRUG OTHER		06/01/2011	

Labeler - Nationwide Laboratories (078366153)