

**CITRANATAL MEDLEY- calcium citrate, iron pentacarbonyl, ferrous fumarate, cholecalciferol, .alpha.-tocopherol, dl-, pyridoxine hydrochloride, folic acid, and doconexent capsule, gelatin coated  
Mission Pharmacal Company**

*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.*

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**CitraNatal Medley**®

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.

Ingestion of more than 3 grams of omega-3 fatty acids per day has been shown to have potential antithrombotic effects, including an 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.

Folic acid alone is improper therapy in the treatment of pernicious anemia and other megaloblastic anemias where vitamin B12 is deficient.

CitraNatal Medley® is a prescription prenatal/postnatal multi-vitamin/mineral soft gel with Ferr-Ease®, a dual-iron delivery comprising both a quick release and slow release iron, and DHA, an essential fatty acid. The prenatal vitamin is an oval, purple, opaque soft gel containing a greenish-gray liquid to semi-solid fill. The prenatal soft gel is printed "596" in white ink.

CitraNatal Medley is a multi-vitamin/mineral prescription drug indicated for use in improving the nutritional status of women prior to conception, throughout pregnancy, and in the postnatal period for both lactating and nonlactating mothers.

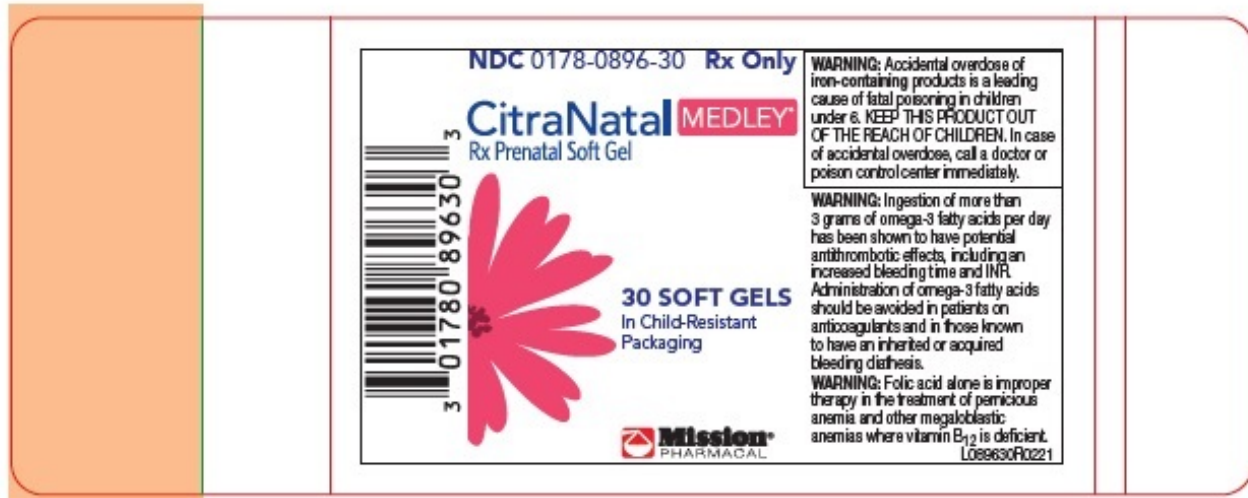
This product is contraindicated in patients with a known hypersensitivity to any of the ingredients.

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

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

One prenatal soft gel daily or as directed by a physician.

Bottles of 30 soft gels each - NDC 00178-0896-30.



## CITRANATAL MEDLEY

calcium citrate, iron pentacarbonyl, ferrous fumarate, cholecalciferol, .alpha.-tocopherol, dl-, pyridoxine hydrochloride, folic acid, and doconexent capsule, gelatin coated

### Product Information

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:0178-0896
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>FERROUS FUMARATE</b> (UNII: R5L488RY0Q) (FERROUS CATION - UNII:GW89581OWR)	FERROUS CATION	2 mg
<b>CALCIUM CITRATE</b> (UNII: MLM29U2X85) (CALCIUM CATION - UNII:2M83C4R6ZB)	CALCIUM CITRATE ANHYDROUS	62 mg
<b>IRON</b> (UNII: E1UOL152H7) (IRON - UNII:E1UOL152H7)	IRON	27 mg
<b>CHOLECALCIFEROL</b> (UNII: 1C6V77QF41) (CHOLECALCIFEROL - UNII:1C6V77QF41)	CHOLECALCIFEROL	200 [IU]

<b>.ALPHA.-TOCOPHEROL</b> (UNII: H4N855PNZ1) (.ALPHA.-TOCOPHEROL - UNII:H4N855PNZ1)	.ALPHA.-TOCOPHEROL	15 [IU]
<b>PYRIDOXINE</b> (UNII: KV2JZ1BI6Z) (PYRIDOXINE - UNII:KV2JZ1BI6Z)	PYRIDOXINE	12.5 mg
<b>FOLIC ACID</b> (UNII: 935E97BOY8) (FOLIC ACID - UNII:935E97BOY8)	FOLIC ACID	1 mg
<b>DOCONEXENT</b> (UNII: ZAD9OKH9JC) (DOCONEXENT - UNII:ZAD9OKH9JC)	DOCONEXENT	200 mg

### Inactive Ingredients

Ingredient Name	Strength
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>GELATIN</b> (UNII: 2G86QN327L)	
<b>LECITHIN, SUNFLOWER</b> (UNII: 834K0WOS5G)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	
<b>BUTYLATED HYDROXYTOLUENE</b> (UNII: 1P9D0Z171K)	
<b>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)	
<b>ETHYL VANILLIN</b> (UNII: YC9ST449YJ)	
<b>D&amp;C RED NO. 33</b> (UNII: 9DBA0SBB0L)	

### Product Characteristics

<b>Color</b>	purple (opaque)	<b>Score</b>	no score
<b>Shape</b>	CAPSULE (Soft Gel)	<b>Size</b>	15mm
<b>Flavor</b>		<b>Imprint Code</b>	596
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0178-0896-30	30 in 1 BOTTLE; Type 0: Not a Combination Product	03/20/2018	



## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		03/20/2018	

**Labeler** - Mission Pharmacal Company (008117095)

**Registrant** - Mission Pharmacal Company (927726893)

## Establishment

Name	Address	ID/FEI	Business Operations
Mission Pharmacal Company		927726893	label(0178-0896) , pack(0178-0896)