SE-NATAL 19- .beta.-carotene, ascorbic acid, cholecalciferol, .alpha.tocopherol acetate, dl-, thiamine, riboflavin, niacinamide, pyridoxine hydrochloride, folic acid, cyanocobalamin, calcium pantothenate, calcium carbonate, ferrous fumarate, zinc oxide and docusate sodium tablet Seton Pharmaceuticals

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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#### Se-Natal 19 Tablets

Vitamins & Minerals with Folic Acid

R<sub>x</sub> Only

### DESCRIPTION

Se-Natal 19 is a white, oblong, film coated tablet, debossed "TL 019". Each film coated tablet contains:

Su	pplemen	t Facts
Servings per Bottle: 100 Serving Size: 1 tablet		
Each Tablet contains:		% DV for Pregnant and Lactating Women
Vitamin A (as beta carotene)	1000 IU	23%
Vitamin C (ascorbic acid)	100 mg	83%
Vitamin D (as cholecalciferol)	10 mcg	67%
Vitamin E (as dl-alpha tocoperol acetate)	30 IU	142%
Thiamine (Vitamin B <sub>1</sub> )	3 mg	214%
Riboflavin (Vitamin B <sub>2</sub> )	3 mg	188%
Niacin (as niacinamide)	15 mg	83%
Vitamin B <sub>6</sub> (as pyridoxine HCl)	20 mg	1000%
Folate (folic acid)	1000 mcg	167%
Vitamin B <sub>12</sub> (as cyanocobalamin)	12 mcg	429%
Pantothenic Acid (as calcium pantothenate)	7 mg	100%
Calcium (as calcium carbonate)	200 mg	15%
lron (as ferrous fumarate)	29 mg	107%
Zinc (as zinc oxide)	20 mg	154%

### **OTHER INGREDIENTS**

Acacia, Croscarmellose Sodium, Fumed Silica, Hypromellose, Magnesium Stearate, Microcrystalline Cellulose, Polyethylene Glycol, Polyvinyl Alcohol, Stearic Acid, Talc and Titanium Dioxide.

# INDICATIONS

Se-Natal 19 tablets are indicated for use in improving the nutritional status of women throughout pregnancy and in the postnatal period for both lactating and non-lactating mothers. Se-Natal 19 tablets are also useful 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.

## WARNINGS

Folic acid alone is improper therapy in the treatment of pernicious anemia and other megaloblastic anemias where vitamin  $B_{12}$  is deficient.

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.

## PRECAUTIONS

### General

Take 2 hours after meals. Do not exceed recommended dose. Discontinue use if symptoms of intolerance appear.

### Folic Acid

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

### **Pediatric Use**

Safety and effectiveness in pediatric patients have not been established.

### Geriatric Use

Dosing for elderly patients should be cautious. Due to the greater frequency of decreased hepatic, renal, or cardiac function, and of concomitant disease or other drug therapy, dosing should start at the lower end of the dosing range.

# **ADVERSE REACTIONS**

Adverse reactions with iron therapy may include GI irritation, constipation, diarrhea, nausea, vomiting, and dark stools. Adverse reactions with iron therapy are usually transient. Allergic sensitization has been reported following both oral and parenteral administration of folic acid.

## DRUG INTERACTIONS

Prescriber should be aware of a number of iron/drug interactions, including antacids, tetracyclines, or fluoroquinolones.

## OVERDOSAGE

### Symptoms:

Abdominal pain, metabolic acidosis, anuria, CNS damage, coma, convulsions, death, dehydration, diffuse vascular congestion, hepatic cirrhosis, hypotension, hypothermia, lethargy, nausea, vomiting, diarrhea, tarry stools, melena, hematemesis, tachycardia, hyperglycemia, drowsiness, pallor, cyanosis, lassitude, seizures, and shock.

## DOSAGE AND ADMINISTRATION

One tablet daily, or as directed by a physician.

## NOTICE

Contact with moisture can discolor or erode the tablet. Do not chew tablet.

## HOW SUPPLIED

Se-Natal 19 tablets for oral administration are supplied as white, coated, oblong tablets debossed "TL019", packaged in bottles of 100 tablets.

### Product Code: 13925-116-01.

### STORAGE

Store at 20° - 25°C (68° - 77°F), excursions permitted to 15° - 30°C (59° - 86°F) [See USP controlled room temperature].

### PHARMACIST

Dispense in a tight, light-resistant container as defined in the USP/NF with child-resistant closure.

Call your doctor about side effects. You may report side effects to FDA at 1-800-FDA-1088.

All prescriptions using this product shall be pursuant to state statutes as applicable. This is not an Orange Book product. This product may be administered only under a physician's supervision. There are no implied or explicit claims on therapeutic equivalence.

### **Reserved for Professional Recommendation.**

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

## R<sub>x</sub> Only

### **Manufactured For:**

Seton Pharmaceuticals Manasquan, NJ 08736 1-800-510-3401

MADE IN CANADA

Code 116-01

Rev. 03/18

# SETON PHARMACEUTICALS

## PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

NDC 13925-**116**-01

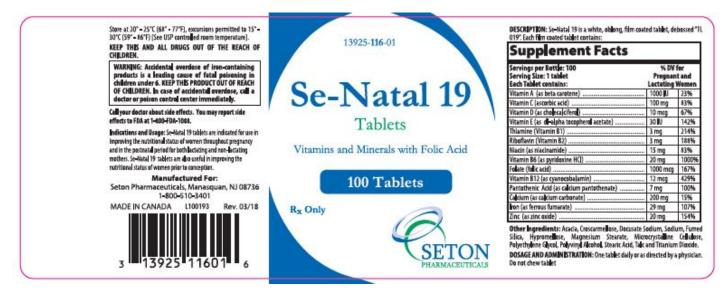
#### Se-Natal 19 Tablets

Vitamins and Minerals with Folic Acid

## 100 Tablets

## R<sub>x</sub> Only

### SETON PHARMACEUTICALS



**SE-NATAL 19** 

.beta.-carotene, ascorbic acid, cholecalciferol, .alpha.-tocopherol acetate, dl-, thiamine, riboflavin, niacinamide, pyridoxine hydrochloride, folic acid, cyanocobalamin, calcium pantothenate, calcium carbonate, ferrous fumarate, zinc oxide and docusate sodium tablet

Product Information					
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (	(Source)	NDC:1	3925-116
Route of Administration	ORAL				
Active Ingredient/Active	Moiety				
Ing	gredient Name		Basis of Streng		Strength
.BETACAROTENE (UNII: 01YAE0	3M7J) (.BETACAROTENE - UNII:01)	AE03M7J)	.BETACAROT	ENE	1000 [iU]
ASCORBIC ACID (UNII: PQ6CK8PD	OR) (ASCORBIC ACID - UNII:PQ6CK	3PD0R)	ASCORBIC AC	ID	100 mg
CHOLECALCIFEROL (UNII: 1C6V7	7QF41) (CHOLECALCIFEROL - UNII:	.C6V77QF41)	CHOLECALCIF	EROL	400 [iU]
<b>.ALPHATOCOPHEROL ACETATI</b> DL UNII:7QWA1RIO01)	E, DL- (UNII: WR1WPI7EW8) (.ALPHA	TOCOPHEROL,	.ALPHA TOCOPHEROL	, DL-	30 mg
THIAMINE (UNII: X66NSO3N35) (TH	HIAMINE ION - UNII:4ABT0J945J)		THIAMINE		3 mg
RIBOFLAVIN (UNII: TLM29760FR) (	RIBOFLAVIN - UNII:TLM2976OFR)		RIBOFLAVIN		3 mg
NIACINAMIDE (UNII: 25X51I8RD4)	(NIACINAMIDE - UNII:25X51I8RD4)		NIACINAMIDE		15 mg
<b>PYRIDOXINE HYDROCHLORIDE</b> ( UNII:KV2JZ1BI6Z)	UNII: 68Y4CF58BV) (PYRIDOXINE -		PYRIDOXINE		20 mg
FOLIC ACID (UNII: 935E97BOY8) (	FOLIC ACID - UNII:935E97BOY8)		FOLIC ACID		1 mg
CYANOCOBALAMIN (UNII: P6YC3	EG204) (CYANOCOBALAMIN - UNII:F	6YC3EG204)	CYANOCOBAL	AMIN	12 ug
CALCIUM PANTOTHENATE (UNII: UNII:19F5HK2737)	568ET80C3D) (PANTOTHENIC ACI	) -	PANTOTHENIC	ACID	7 mg
CALCIUM CARBONATE (UNII: HOC	69379FGK) (CALCIUM CATION - UNI	:2M83C4R6ZB)	CALCIUM CATI	ON	200 mg
FERROUS FUMARATE (UNII: R5L4	88RY0Q) (FERROUS CATION - UNII:	GW895810WR)	FERROUS CAT	ION	29 mg
ZINC OXIDE (UNII: SOI2LOH54Z) (	ZINC CATION - UNII:13S1S8SF37)		ZINC CATION		20 mg
DOCUSATE SODIUM (UNII: F05Q2	T2JA0) (DOCUSATE - UNII:M7P2719	5AG)	DOCUSATE SO	DIUM	25 mg
Inactive Ingredients					
	Ingredient Name			St	rength
ACACIA (UNII: 5C5403N26O)					
CROSCARMELLOSE SODIUM (UN	III: M28OL1HH48)				
SILICON DIOXIDE (UNII: ETJ7Z6XE	3U4)				
HYPROMELLOSES (UNII: 3NXW29)	/3WO)				
MAGNESIUM STEARATE (UNII: 70	097M6I30)				
CELLULOSE, MICROCRYSTALLIN	IE (UNII: OP1R32D61U)				

POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)

POLYVINYL ALCOHOL (UNII: 532B59J990)

**STEARIC ACID** (UNII: 4ELV7Z65AP)

TALC (UNII: 7SEV7J4R1U)

TITANIUM DIOXIDE (UNII: 15FIX9V2JP)

	-		
Product Characterist	tics		
Color	WHITE	Score	no score

5	аре		CAPSULE	Size		19mm
Fla	vor			Imprint Cod	e	TL019
Co	ntains					
Pa	ackaging					
#	Item Code		Package Descript	tion	Marketing Start Date	Marketing End Date
_	NDC:13925-116-	100 in 1	BOTTLE; Type 0: Not a (	Combination	07/01/2000	
	01	Product			07/01/2009	
					07/01/2009	
•	01	Product			07/01/2009	
•		Product	nation		07/01/2009	
±	01	Product	<b>nation</b> plication Number or Citation		Marketing Start Date	Marketing End Date

Labeler - Seton Pharmaceuticals (828898002)

Registrant - Seton Pharmaceuticals (828898002)

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Seton Pharmaceuticals