

METAVEX- metavex tablet

Oncora Pharma, LLC

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.

Metavex

DESCRIPTION

Metavex™ is a prescription folic acid-containing dietary supplement intended to support metabolic health. Each tablet contains a combination of vitamins, minerals, and botanical ingredients formulated to provide nutritional support.

INDICATIONS AND USAGE

Metavex™ is indicated as a prescription dietary supplement for the dietary management of individuals with metabolic health needs where folic acid supplementation is appropriate.

Administration:

Take two (2) tablets daily with a 12-ounce glass of water, or as directed by a licensed healthcare practitioner.

CONTRAINDICATIONS

Metavex™ is contraindicated in patients with known hypersensitivity to any of its components.

WARNINGS

Metavex™ should be administered under the supervision of a licensed medical practitioner.

Patients should not exceed the recommended dosage.

If adverse reactions occur, discontinue use and consult a healthcare professional.

Precautions

Folic Acid and Vitamin B12 Deficiency:

Folic acid administered alone may be inappropriate for the treatment of pernicious anemia or other megaloblastic anemias where vitamin B12 deficiency is present. Doses of folic acid greater than 0.1 mg daily may mask hematologic signs of pernicious anemia while neurological manifestations continue to progress.

Patients with underlying medical conditions or those taking other medications should consult their healthcare provider prior to use.

DOSAGE AND ADMINISTRATION

Take two (2) tablets daily with a 12-ounce glass of water, or as directed by a licensed healthcare practitioner.

HOW SUPPLIED

Metavex™ is supplied as tan, oval tablets in bottles of 30 tablets.

NDC 85477-503-30

Display Panel

**METAVEX™ SHOULD BE
ADMINISTERED UNDER THE
SUPERVISION OF A LICENSED
MEDICAL PRACTITIONER.**

DOSAGE AND ADMINISTRATION:
Usual adult dose is 2 tablets once or twice daily taken with 8-12 oz of water or as prescribed by a licensed practitioner.

**KEEP OUT OF REACH OF CHILDREN.
CALL A DOCTOR OR POISON
CONTROL CENTER IMMEDIATELY IN
CASE OF ACCIDENTAL OVERDOSE.**
Call your licensed medical practitioner about side effects. You may report side effects by calling Oncora Pharma at 1-800-541-4802 or 1-800-FDA-1088.

STORAGE: Store at 20°-25°C (68°-77°F); excursions permitted to 15°-30°C (59°-86°F). Protect from light and moisture.

DISTRIBUTED BY:
Oncora Pharma Dallas, TX 75228

85477-503-30

Metavex™

METABOLIC HEALTH

Prescription Folic Acid-Containing Dietary Supplement

30 TABLETS

Dispensed by Prescription

SUPPLEMENTAL FACTS

Amount Per Serving:	% Daily Value
Vitamin D (as cholecalciferol) 250 mcg (10,000 IU)	1250%
Folate (from 2000 mcg folic acid) 3,400 mcg DFE	850%
Inulin (from organic agave (heart) extract) 1,000 mg *	*
Slippery Elm (bark) 850 mg *	*
Aloe Vera (leaf) 450 mg *	*
Cascara sagrada (bark) 270 mg *	*

*Daily value not established

OTHER INGREDIENTS: Stearic Acid, Calcium Carbonate, Di-Calcium Phosphate, Magnesium Stearate

©2025 Oncora Pharma

3 85477 00003 7

Metavex™

METABOLIC HEALTH

Prescription Folic Acid-Containing Dietary Supplement

30 Tablets

Dispensed by Prescription

Distributed by:
Oncora Pharma
Dallas, TX 75228

NDC **85477-503-30**

METAVEX

metavex tablet

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:85477-503
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHOLECALCIFEROL (UNII: 1C6V77QF41) (CHOLECALCIFEROL - UNII:1C6V77QF41)	CHOLECALCIFEROL	.125 mg
FOLIC ACID (UNII: 935E97BOY8) (FOLIC ACID - UNII:935E97BOY8)	FOLIC ACID	1 mg
BLUE AGAVE INULIN (AGAVE TEQUILANA) (UNII: PW973NF38H) (BLUE AGAVE INULIN (AGAVE TEQUILANA) - UNII:PW973NF38H)	BLUE AGAVE INULIN (AGAVE TEQUILANA)	500 mg
ELM (UNII: 63POE2M46Y) (ELM - UNII:63POE2M46Y)	ELM	425 mg
ALOE FEROX LEAF (UNII: 0D145J8EME) (ALOE FEROX LEAF - UNII:0D145J8EME)	ALOE FEROX LEAF	225 mg
RHAMNUS CATHARTICA BARK (UNII: KD27950XHY) (RHAMNUS CATHARTICA BARK - UNII:KD27950XHY)	RHAMNUS CATHARTICA BARK	135 mg

Inactive Ingredients

Ingredient Name	Strength
STEARIC ACID (UNII: 4ELV7Z65AP)	
CALCIUM CARBONATE (UNII: H0G9379FGK)	
CALCIUM PHOSPHATE (UNII: 97Z1W3NDX)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	

Product Characteristics

Color	brown	Score	no score
Shape	OVAL	Size	23mm
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:85477-503-30	1 in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		01/01/2026	

Labeler - Oncora Pharma, LLC (119482542)

