

**VIRELIX- virelix tablet**  
**Oncora Pharma, LLC**

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**Virelix**

**Health Claim Section:**

Virelix Multivitamin

Dispensed by Prescription

**Description:**

85622-533-30

Reserved for Professional Recommendation

Virelix Tablets should be administered under the supervision of a licensed medical practitioner.

Virelix Tablets are orally administered prescription dietary supplement tablets intended to provide significant amounts of Vitamins B1, B3, B6, B12, C, D3, folic acid, and NADH to supplement the diet and help assure that nutritional deficiencies of these vitamins will not develop.

This listed product is not a National Drug Code, but instead has merely been formatted to comply with standard industry practice for pharmacy and insurance computer systems.

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.

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.

Manufactured for:

Oncora Pharma

Dallas TX 75228

**Warnings:**

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

Virelix Tablets should only be used under the direction and supervision of a licensed medical practitioner. Use with caution in patients that may have a medical condition, are pregnant, lactating, trying to conceive, under the age of 18, or taking medications.

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.

**Dosage and Administration:**

Adults: One tablet daily as a dietary supplement, preferably with a meal, or as recommended by a healthcare professional.

**Precautions:**

## CONTRAINDICATIONS

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

## PRECAUTIONS

Use under the direction and supervision of a licensed healthcare professional.

## ADVERSE REACTIONS

Allergic sensitization has been reported following oral administration of folic acid. Consult your physician immediately if adverse reactions occur.

KEEP OUT OF THE REACH OF CHILDREN.

**Storage:**

Store at 20° to 25°C (68° to 77°F); excursions permitted to 15° to 30°C (59° to 86°F).

**Label:**

**DOSAGE AND ADMINISTRATION:**

Usual adult dosage is 1 tablet taken orally once daily or as prescribed by a licensed medical practitioner.

**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.

**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.

NDC 85477-533-30 Rx Only

# VIRELIX

DIETARY SUPPLEMENT

Dispensed by Prescription

30 Tablets

**SUPPLEMENTAL FACTS**

Amount Per Serving:	% Daily Value
Vitamin C (as ascorbic acid)	125 mg 139%
Vitamin D3 (as cholecalciferol)	12.5 mcg 63%
Vitamin B1 (as Thiamin HCl)	25 mg 2083%
Vitamin B6 (as pyridoxal phosphate anhydrous)	125 mg 735%
Folic Acid	1.667 mg 417%
Vitamin B12 (as methylcobalamin)	1 mg 41,677%
NADH (reduced nicotinamide adenine dinucleotide)	5 mg †
CoEnzyme Q-10 (ubiquinone)	50 mg †

† Daily Value (DV) not established.

**Inactive Ingredients:** Microcrystalline Cellulose, Pregelatinized Starch, Coating (Beta-Carotene, Titanium Dioxide, Hypromellose, Polyvinyl Alcohol, Polyethylene Glycol, Talc), Croscarmellose Sodium, Silica, Silicon Dioxide, Hydroxypropyl Cellulose, Magnesium Stearate.



Distributed by:  
Oncora Pharma  
Dallas, TX 75228  
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LOT:      EXP:

pdp

## VIRELIX

virelix tablet

### Product Information

<b>Product Type</b>	DIETARY SUPPLEMENT	<b>Item Code (Source)</b>	NHRIC:85477-533
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>ASCORBIC ACID</b> (UNII: PQ6CK8PD0R) (ASCORBIC ACID - UNII:PQ6CK8PD0R)	ASCORBIC ACID	125 mg
<b>CHOLECALCIFEROL</b> (UNII: 1C6V77QF41) (CHOLECALCIFEROL - UNII:1C6V77QF41)	CHOLECALCIFEROL	.125 mg
<b>THIAMINE HYDROCHLORIDE</b> (UNII: M572600E5P) (Thiamine ION - UNII:4ABT0J945J)	THIAMINE HYDROCHLORIDE	25 mg
<b>PYRIDOXINE HYDROCHLORIDE</b> (UNII: 68Y4CF58BV) (PYRIDOXINE - UNII:KV2JZ1BI6Z)	PYRIDOXINE HYDROCHLORIDE	125 mg
<b>FOLIC ACID</b> (UNII: 935E97BOY8) (FOLIC ACID - UNII:935E97BOY8)	FOLIC ACID	1.667 mg

<b>CYANOCOBALAMIN</b> (UNII: P6YC3EG204) (CYANOCOBALAMIN - UNII:P6YC3EG204)	CYANOCOBALAMIN	1 mg
<b>NICOTINAMIDE ADENINE DINUCLEOTIDE</b> (UNII: 0U46U6E8UK) (NICOTINAMIDE ADENINE DINUCLEOTIDE - UNII:0U46U6E8UK)	NICOTINAMIDE ADENINE DINUCLEOTIDE	5 mg
<b>UBIQUINONE</b> (UNII: EJ27X76M46) (UBIQUINONE - UNII:EJ27X76M46)	UBIQUINONE	50 mg

### Inactive Ingredients

Ingredient Name	Strength
<b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)	
<b>CROSCARMELLOSE SODIUM</b> (UNII: M28OL1HH48)	
<b>SILICA</b> (UNII: ETJ7Z6XBU4)	
<b>HYDROXYPROPYL CELLULOSE (110000 WAMW)</b> (UNII: 5Y0974F5PW)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NHRIC:85477-533-30	30 in 1 PACKAGE		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
DIETARY SUPPLEMENT		10/09/2025	

### Supplement Facts

Serving Size :		Serving per Container :	
	Amount Per Serving		% Daily Value
color			
scoring	1		
shape			
size (solid drugs)	18 mm		

**Labeler** - Oncora Pharma, LLC (119482542)