NEPHRON FA- mineral/vitamin supplement tablet, coated
Nephro-Tech, Inc.

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.

Nephron FA
multi-vitamin/mineral supplement with ascorbic acid, thiamine mononitrate, riboflavin, niacinamide, pyridoxine hydrochloride, folic acid, cyanocabalimin, biotin, pantothenic acid and ferrous fumarate, film-coated

Description

Nephron FA is a prescription folic acid supplement with additional nutrients for kidney dialysis patients. Nephron FA is an elongated, white, film coated tablet imprinted with NT and scored on one side, plain on the other.

Precaution

Folic Acid may obscure pernicious anemia or produce remission while neurologic progress may continue.

Suggested Dosage

One tablet daily taken away from meals, or as directed by the physician.

Package Label/Principal Display Panel

Nephron FA label

59528-4456-1

Rx Only

Nephron FA

Vitamin/Mineral Supplement

100 tablets


100 Tablets
59528-4456-1
NEPHRON FA®
Vitamin/Mineral Supplement
℞ only
Dist. by
NEPHRO-TECH, INC.
P.O. Box 16106
Shawnee, Kansas 66203
SUPPLEMENT FACTS
Serving Size 1 Tablet
Amount Per Tablet % Daily Value
Vitamin C (as ascorbic acid) 40 mg 67%
Thiamin (as thiamine mononitrate) 1.5 mg 100%
Riboflavin 1.7mg 100%
Niacin (as niacinamide) 16 mg NE 100%
Vitamin B6 (as pyridoxine hydrochloride) 10 mg 500%
Folate 1,700 mcg DFE 425%
(1,000 mcg folic acid)
Vitamin B12 (as cobalamin) 6 mcg 100%
Biotin 300 mcg 100%
Pantothenic Acid (as d-calcium pantothenate) 10 mg 100%
Iron (as ferrous fumarate) 66 mg 367%
*Percent Daily Value based on 2,000 calorie diet.
Other Ingredients: microcrystalline cellulose, hydrogenated vegetable oil, docusate sodium, stearic acid, sodium croscarmellose, dibasic calcium phosphate, sodium benzoate, magnesium stearate, hydroxypropylmethylcellulose, silicon dioxide, titanium dioxide, maltodextrin, triacetin, hydroxypropylcellulose, polyethylene glycol, sodium citrate, resin.
Precaution: Folic Acid may obscure pernicious anemia or produce remission while neurologic progress may continue.
Suggested Dosage: One tablet daily taken away from meals, or as directed by the physician.
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.
Tamper Evident: Do not use if printed inner seal is broken.
Rev. 11/19 QL-00142

NEPHRON FA

mineral/vitamin supplement tablet, coated

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:59528-4456
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
IRON (UNII: E1UOL152H7) (FERROUS CATION - UNII:GW89581OWR)	FERROUS CATION	66 mg
ASCORBIC ACID (UNII: PQ6CK8PD0R) (ASCORBIC ACID - UNII:PQ6CK8PD0R)	ASCORBIC ACID	40 mg
NIACINAMIDE (UNII: 25X51I8RD4) (NIACINAMIDE - UNII:25X51I8RD4)	NIACINAMIDE	20 mg
PYRIDOXINE HYDROCHLORIDE (UNII: 68Y4CF58BV) (PYRIDOXINE - UNII:KV2JZ1BI6Z)	PYRIDOXINE HYDROCHLORIDE	10 mg
PANTOTHENIC ACID (UNII: 19F5HK2737) (PANTOTHENIC ACID - UNII:19F5HK2737)	PANTOTHENIC ACID	10 mg
RIBOFLAVIN (UNII: TLM2976OFR) (RIBOFLAVIN - UNII:TLM2976OFR)	RIBOFLAVIN	1.7 mg
THIAMINE (UNII: X66NSO3N35) (THIAMINE ION - UNII:4ABT0J945J)	THIAMINE	1.5 mg
FOLIC ACID (UNII: 935E97BOY8) (FOLIC ACID - UNII:935E97BOY8)	FOLIC ACID	1 mg
Biotin (UNII: 6SO6U10H04) (BIOTIN - UNII:6SO6U10H04)	Biotin	.3 mg
COBALAMIN (UNII: 8406EY2OQA) (COBALAMIN - UNII:8406EY2OQA)	COBALAMIN	.006 mg

Product Characteristics

Color	white	**Score**	2 pieces
Shape	capsule	**Size**	19mm
Flavor		**Imprint Code**	NT
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59528-4456-1	100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/01/1992	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		11/01/1992	

Labeler - Nephro-Tech, Inc. (878520485)

Registrant - Nephro-Tech, Inc. (878520485)

Establishment

Name	Address	ID/FEI	Business Operations
LGM Pharma		117549200	manufacture(59528-4456)

Revised: 7/2021

Nephro-Tech, Inc.