DAVIMET WITH IRON- iron, folate tablet PureTek Corporation

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.

Davimet with Iron

DESCRIPTION:

Each caplet contains: Vitamin A (as Retinyl Acetate)750 mcg RAE	
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Vitamin C (as Sodium Ascorbate 36 mg and Ascorbic Acid 24 mg) 60 mg	1
Vitamin D3 (as Cholecalciferol) 10 mcg (400 IU)	
Vitamin E (as DL-Alpha Tocopheryl Acetate) 6.75 mg	
Thiamin (as Thiamine Mononitrate) 1.05 mg	
Riboflavin 1.2 mg	
Niacin (as Niacinamide) 13.5 mg	
Vitamin B6 (as Pyridoxine Hydrochloride) 1.05 mg	
Folate (as L-5-Methyltetrahydrofolate calcium salt)1700 mcg DFE	
(1000 mcg as L-5-Methylfolate)	
Vitamin B12 (as Methylcobalamin)	
Iron (as Iron Amino Acid Chelate) 11 mg	

Other Ingredients: Aspartame, CI 15985 / [Yellow 6 Lake], Croscarmellose Sodium, Magnesium Stearate, Maltodextrin, Microcrystalline Cellulose, Natural OrangeFlavor, Stearic Acid (Vegetable Source), Sucrose.

Phenylketonurics: Phenylalanine 3.0 mg Per Tablet.

INDICATIONS AND USAGE:

Davimet™ with Iron Multivitamin Chewable Tablets are indicated for the treatment of iron deficiency anemia and folate deficiency, commonly seen in conditions such as extended convalescence, menorrhagia, pregnancy, puberty, excessive blood loss, and advanced age. They are also indicated for condition in which irondeficiency and vitamin C deficiency occur together, along with a deficient intake or increased need for B-Complex vitamins in chronic and acute illness, as well as cases of metabolic stress, and in convalescence.

CONTRAINDICATIONS:

This product is contraindicated in patients with known hypersensitivity to any of its ingredients; also, all iron compounds are contraindicated in patients withhemosiderosis, hemochromatosis, or hemolytic anemias. Pernicious anemia is a contraindication, as folic acid may obscure its signs and symptoms.

WARNING:

Keep out of the reach of children. In case of accidental overdose, seek professional assistance or contact a Poison Control Center immediately.

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 accidentaloverdose, call a doctor or poison control center immediately.

Administration of folate alone is improper therapy for pernicious anemia and other megaloblastic anemias in which vitamin B12 is deficient.

CAUTION: Should be chewed.

PRECAUTIONS:

Folate in doses above 0.1 mg daily may obscure pernicious anemia, in that hematologic remission can occur while neurological manifestations remain progressive. There is a potential danger in administering folate to patients with undiagnosed anemia, since folate may obscure the diagnosis of pernicious anemia by alleviating the hematologic manifestations of the disease while allowing the neurologic complications to progress. This may result in severe nervous system damage before the correct diagnosis is made. Adequate doses of vitamin B12 may prevent, halt, or improve the neurologic changes caused by pernicious anemia.

The patient's medical conditions and consumption of other drugs, herbs, and/or supplements should be considered.

Use only under the direction of a licensed healthcare practitioner. Call your doctor about side effects. To report side effects, call **PureTek Corporation** at **1-877-921-7873** or **FDA** at **1-800-FDA-1088** or **www.fda.gov/medwatch.**

Drug Interactions:

Davimet™ with Iron Multivitamin Chewable Tablets are not recommended and should not be given to patients receiving levodopa because the action oflevodopa is antagonized by pyridoxine. There is a possibility of increased bleeding due to pyridoxine interaction with anticoagulants (e.g., Aspirin, Heparin orClopidogrel).

Adverse Reactions:

Folate: Allergic sensitizations have been reported following both oral and parenteral administration of folate.

Iron: Gastrointestinal disturbances (anorexia, nausea, diarrhea, constipation) occur occasionally, but are usually mild and may subside with continuation of therapy. Although the absorption of iron is best when taken between meals, giving **Davimet™ with Iron Multivitamin Chewable Tablets** after meals may controloccasional gastrointestinal disturbances. **Davimet™ with Iron Multivitamin Chewable Tablets** is best absorbed when taken at bedtime.

Adverse reactions have been reported with specific vitamins and minerals but generally

at levels substantially higher than those contained herein. However, allergicand idiosyncratic reactions are possible at lower levels. Iron, even at the usual recommended levels, has been associated with gastrointestinal intolerance in somepatients

OVERDOSE:

Iron: Signs and Symptoms: Iron is toxic. Acute overdosage of iron may cause nausea and vomiting and, in severe cases, cardiovascular collapse and death. Othersymptoms include pallor and cyanosis, melena, shock, drowsiness and coma. The estimated overdose of orally ingested iron is 300 mg/kg body weight. Whenoverdoses are ingested by children, severe reactions, including fatalities, have resulted. **Davimet™ with Iron Multivitamin Chewable Tablets** should be stored beyond the reach of children to prevent against accidental iron poisoning.

Treatment:

For specific therapy, exchange transfusion and chelating agents should be used. For general management, perform gastric lavage with sodium bicarbonate solutionor milk. Administer intravenous fluids and electrolytes and use oxygen.

DOSAGE AND ADMINISTRATION:

One tablet daily, to be dissolved in the mouth or chewed before swallowing.

Do not use if bottle seal is broken.

KEEP THIS AND ALL MEDICATIONS OUT OF THE REACH OF CHILDREN.

Store at 20° to 25°C (68° to 77°F) [see USP Controlled Room Temperature].

HOW SUPPLIED:

Davimet™ with Iron Multivitamin Chewable Tablets are light orange with speckles, orange flavor, un-scored, round tablets. Available on prescription only in bottleof 30 tablets – NDC 59088-160-54. Dispense in a tight, light resistant container with a child resistant closure as defined in the USP/NF. All prescription substitutions using this product shall be pursuant to state statutes as applicable. This is not an Orange Book product.

Davimet™ with Iron

Manufactured by: PureTek Corporation Panorama City, CA 91402

For questions or information call toll-free: **877-921-7873**



Supplement Facts

Serving Size: 1 Tablet Serving per container: 30

	Amount Per Serving %	Daily Value
Vitamin A (as Retinyl Acetate)	750 mcg RAE	83%
Vitamin C (as Sodium Ascorbate 36 mg and Ascorbi	c Acid 24 mg) 60 mg	67%
Vitamin D3 (as Cholecalciferol)	10 mcg (400 IU)	50%
Vitamin E (as DL-Alpha Tocopheryl Acetate)	6.75 mg	45%
Thiamin (as Thiamine Mononitrate)	1.05 mg	88%
Riboflavin	1.2 mg	92%
Niacin (as Niacinamide)	13.5 mg	84%
Vitamin Ba (as Pyridoxine Hydrochoride)	1.05 mg	62%
Folate (as L-5-Methyttetrahydrofolate calcium saft) (1000 mc)	1700 mcg DFE g of L-5-methylfolate)	425%
Vitamin B12 (as Methylcobalamin)	4.5 mcg	188%
Iron (as Iron Amino Acid Chelate)	11 mg	61%

NDC 59088-160-54 Rx Only

DERMACINE

Davimet™

with Iron

Multivitamin Chewable Tablets

Orange Flavor

30 Tablets

DAVIMET WITH IRON

iron, folate tablet

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:59088-160
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
VITAMIN A ACETATE (UNII: 3LE3D9D6OY) (VITAMIN A - UNII:81G40H8B0T)	VITAMIN A	750 ug
ASCORBIC ACID (UNII: PQ6CK8PD0R) (ASCORBIC ACID - UNII:PQ6CK8PD0R)	ASCORBIC ACID	24 mg
NIACINAMIDE (UNII: 25X5118RD4) (NIACINAMIDE - UNII:25X5118RD4)	NIACINAMIDE	13.5 mg
PYRIDOXINE HYDROCHLORIDE (UNII: 68Y4CF58BV) (PYRIDOXINE - UNII: KV2JZ 1BI6Z)	PYRIDOXINE	1.05 mg
THIAMINE MONONITRATE (UNII: 8K0I04919X) (THIAMINE ION - UNII:4ABT0J945J)	THIAMINE	1.05 mg
CHOLECALCIFEROL (UNII: 1C6V77QF41) (CHOLECALCIFEROL - UNII:1C6V77QF41)	CHOLECALCIFEROL	10 ug
IRON (UNII: E1UOL152H7) (IRON - UNII:E1UOL152H7)	IRON	11 mg
RIBOFLAVIN (UNII: TLM29760FR) (RIBOFLAVIN - UNII:TLM29760FR)	RIBOFLAVIN	1.2 mg
METHYLCOBALAMIN (UNII: BR1SN1JS2W) (METHYLCOBALAMIN - UNII:BR1SN1JS2W)	METHYLCOBALAMIN	4.5 ug
SODIUM ASCORBATE (UNII: S033EH8359) (ASCORBIC ACID - UNII:PQ6CK8PD0R)	ASCORBIC ACID	36 mg
LEVOMEFOLATE CALCIUM (UNII: A9R10K3F2F) (LEVOMEFOLIC ACID - UNII:8S95DH25XC)	LEVOMEFOLATE CALCIUM	1000 ug
.ALPHATOCOPHEROL ACETATE, DL- (UNII: WR1WPI7EW8) (.ALPHATOCOPHEROL, DL UNII:7QWA1RIO01)	.ALPHA TOCOPHEROL, DL-	6.75 mg

Inactive Ingredients				
Ingredient Name	Strength			
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)				
MALTODEXTRIN (UNII: 7CVR7L4A2D)				
MAGNESIUM STEARATE (UNII: 70097M6I30)				
ASPARTAME (UNII: Z0H242BBR1)				
SUCROSE (UNII: C151H8M554)				

FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
CROSCARMELLOSE SODIUM (UNII: M280L1HH48)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Product Characteristics				
Color	orange (Light Orange with Speckles)	Score	no score	
Shape	ROUND	Size	13mm	
Flavor		Imprint Code		
Contains				

l	P	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
	1	NDC:59088- 160-54	30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/03/2024		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		06/03/2024	

Labeler - PureTek Corporation (785961046)

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