

FERROUS SULFATE- ferrous sulfate tablet
Richmond Pharmaceuticals, 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.

FERROUS SULFATE TABLETS 325 mg

Supplement Facts[/S]

Serving Size: 1 Tablet	% Daily Value
Amount per Tablet	
Iron (as ferrous sulfate) 65 mg	360 %

SUGGESTED USE

Adults:One (1) tablet daily, preferably after meals or as directed by the doctor.

Children:As directed by the doctor.

Active Ingredient

EACH TABLET CONTAINS: %U.S. RDA*

Elemental Iron 65 mg 360

(Equivalent to 325 mg of Ferrous Sulfate)

* U.S. Recommended Daily Allowance

Inactive Ingredients

Croscarmellose sodium, dicalcium phosphate, FD&C RED#40 (Al-lake), FD&C yellow #6 (Al-lake), hypromellose, magnesium stearate, microcrystalline cellulose, PEG 400, titanium dioxide

Purpose

Iron Supplement

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.

WARNINGS: Do not exceed recommended dosage.The treatment of any anemic

condition should be under the advice and supervision of doctor. Occasional gastrointestinal discomfort (such as nausea) may be minimized by taking with meals. Iron-containing medication may occasionally cause constipation or diarrhea.

As with any drug, if you are pregnant or nursing baby, seek the advice of a health professional before using this product.

DRUG INTERACTION PRECAUTION

Since oral iron products interfere with absorption of oral tetracycline antibiotics , these products should not be taken within two hours of each other.

DOSAGE AND ADMINISTRATION

Each tablet contains:Calcium 20 mg (2% daily value) Store in a dry place at controlled room temperature at 15-30 °C (59°-86° F). Do not expose to excessive heat or moisture.

Questions or Comments

DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS MISSING OR DAMAGED

Call 804-270-4498 Monday-Friday, 8.30 am - 4.30 pm ET

Package Label

FERROUS SULFATE TABLETS 325 mg (5 gr)

Iron Supplement

NDC: 54738-963-01 - 100 Tablets

NDC: 54738-963-03 - 1000 TABLETS

*Compare to Active Ingredient in Feosol®

Ferrous Sulfate

325 mg

IRON SUPPLEMENT

100 TABLETS

**Richmond
Pharmaceuticals, Inc.**

Supplement Facts
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Elemental Iron 65 mg 360
(Equivalent to 325 mg of Ferrous Sulfate)
*U.S. Recommended Daily Allowance

Other ingredients: croscarmellose sodium, dicalcium phosphate, FD&C red #40 (Al-lake), FD&C yellow #6 (Al-lake), hypromellose, magnesium stearate, microcrystalline cellulose, PEG 400, titanium dioxide

TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

*Richmond Pharmaceuticals Inc. is not affiliated with the owner of the registered trademark Feosol®

Distributed by: Richmond Pharmaceuticals, Inc.
Richmond, VA 23233, USA

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Questions or comments:
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FERROUS SULFATE

ferrous sulfate tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54738-963
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
FERROUS SULFATE (UNII: 39R4TAN1VT) (FERROUS CATION - UNII:GW89581OWR)	FERROUS CATION	325 mg

Inactive Ingredients

Ingredient Name	Strength
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	

POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)

TITANIUM DIOXIDE (UNII: 15FIX9V2JP)

Product Characteristics

Color	red	Score	no score
Shape	ROUND	Size	8mm
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54738-963-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	06/01/2015	
2	NDC:54738-963-03	1000 in 1 BOTTLE; Type 0: Not a Combination Product	06/01/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		06/01/2015	

Labeler - Richmond Pharmaceuticals, Inc. (043569607)

Registrant - Advance Pharmaceutical Inc. (078301063)

Establishment

Name	Address	ID/FEI	Business Operations
Advance Pharmaceutical Inc.		078301063	manufacture(54738-963)

Revised: 12/2024

Richmond Pharmaceuticals, Inc.