FERROUS GLUCONATE- ferrous gluconate tablet
Nationwide Pharmaceutical LLC

Ferrous Gluconate Tablets

CONTAINS

Each tablet contains 324 mg of ferrous gluconate, equivalant to 38 mg of elemental iron, providing 208% of the U.S. recommended daily intake (RDI) of iron for adults and children 4 and older.

WARNINGS

Do not take within 2 hours of taking oral tetracycline antibiotics, since oral iron products tend to interfere with absorption of tetracycline. May cause grastrointestinal discomfort, nausea, constipation or diarrhea. If you are pregnant or nursing a baby, seek the advice of a health professional before using this product. U.S. Consumer Product Safety Commission requires that iron-containing medicines and vitamins with iron be packaged in a child-resistant closure. Parents should always properly resecure safety closures.

DIRECTIONS FOR USE

Do not crush or chew tablets.

Adults Serving Size: 1 tablet three to four times daily.
Children: Consult a physician

STORAGE

Store at 15-30°C (58-86°F)

Supplement Facts **Serving Size: 1 Tablet**		
Amount Per Tablet		**% Daily Value**
Iron (as Ferrous Gluconate)	38 mg	208%

Other ingredients: Microcrystalline Cellulose, Sodium Starch Glycolate, Corn Starch, Colloidal Silicon Dioxide, Stearic Acid, Magnesium Stearate, Titanium Dioxide, Polyethylene Glycol, FD&C Blue No.1, Hypromellose, Carnauba Wax.

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.

Manufactured for/Distributed by:
Nationwide Pharmaceutical LLC
San Antonio, TX 78216
Rev 10/2024

PRINCIPAL DISPLAY PANEL - 324 mg Tablet Bottle Label

Nationwide
Pharmaceutical

NDC 69375-0009-10

Ferrous Gluconate
Tablets

324 mg

Dietary Supplement

100 TABLETS

Made in the USA

FERROUS GLUCONATE

ferrous gluconate tablet

Product Information			
Product Type	DIETARY SUPPLEMENT	**Item Code (Source)**	NHRIC:69375-009
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
FERROUS GLUCONATE (UNII: U1B11I423Z) (FERROUS CATION - UNII:GW89581OWR)	FERROUS CATION	38 mg

Inactive Ingredients	
Ingredient Name	**Strength**
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
SODIUM STARCH GLYCOLATE Type B Potato (UNII: 27NA468985)	
STARCH, CORN (UNII: O8232NY3SJ)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
CARNAUBA WAX (UNII: R12CBM0EIZ)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NHRIC:69375-009-10	100 in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
Dietary Supplement		10/01/2024	

Supplement Facts

Serving Size : **Serving per Container :**

	Amount Per Serving	% Daily Value
color		
scoring	1	
shape		
size (solid drugs)	11 mm	

Labeler - Nationwide Pharmaceutical LLC (079265801)

Revised: 10/2024 Nationwide Pharmaceutical LLC