FUSION PLUS- iron / folic acid / vitamin / probiotic supplement capsule
U.S. Pharmaceutical Corporation

FUSION Plus™

Supplement Facts Serving Size: 1 Capsule			
Amount Per Serving		%DV*	% Daily Value for Pregnant or Lactating Women
Vitamin C (as Sodium Ascorbate)	75 mg	83%	63%
Vitamin B_1 (as Thiamine Hydrochloride)	2 mg	167%	143%
Vitamin B_2 (Riboflavin)	3 mg	231%	188%
Vitamin B_3 (as Niacinamide)	10 mg	63%	56%
Vitamin B_6 (Pyridoxine HCl)	10 mg	588%	500%
Folic Acid	1.25 mg	313%	208%
Vitamin B_{12} (Cyanocobalamin)	12 mcg	500%	429%
Biotin	300 mcg	1000%	857%
Pantothenic Acid (as Calcium D-Pantothenate)	6 mg	120%	86%
Iron (65 mg from Ferrous Fumarate) (65 mg from Polysaccharide Iron Complex)	130 mg	722%	481%
Lactobacillus casei (KE-99) 200 Billion CFU/g**	30 mg	†	†
* % Daily Values are based on a 2,000 calorie intake of adults and children 12 years and older. † Daily value not established **Colony Forming Units (CFU) per gram at time of manufacturing			

Other Ingredients: Titanium Dioxide, Candurin Silver Fine, Carmine, Magnesium Stearate, Microcrystalline Cellulose and Hypromellose.

Patent Numbers: USA:6,797,266;5,626,883;Mexico MX/a/2008/004461; Singapore: 200802623-9 and other countries.

INDICATIONS: Fusion Plus™ is indicated for the treatment of iron deficiency anemia

and folate deficiency as in extended convalescence, menorrhagia, pregnancy, puberty, excessive blood loss and advanced age. Also for treatment of condition in which iron deficiency 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: Fusion Plus™ is contraindicated in patients with a known hypersensitivity to any of its ingredients; also, all iron compounds are contraindicated in patients with hemosiderosis, hemochromatosis, or hemolytic anemias. It is also contraindicated in patients suffering from pernicious anemia as folic acid may obscure its signs and symptoms.

Ferrous Fumarate and Polysaccharide Iron Complex: All Fusion™ products include a unique patented source of iron, e.g. Ferrous Fumarate and Polysaccharide Iron Complex (U.S. Patent No: 11/243,043 Pending). An increase in tolerability is observed with the (patented formulation) and is believed to occur as the result of distributing the total iron content in the composition among compounds that provide iron to the patients blood stream via two different mechanisms. The ferrous salts are readily absorbed in the upper gut, by direct dissolution and absorption of the ferrous iron by the bloodstream. However, the iron available from PIC is absorbed in the lower gut, via an active protein transport mechanism.

Clinical Studies: Picinni, L.-Ricciotti, M. 1982. Therapeutic effectiveness of an iron-polysaccharide complex in comparison with iron fumarate in the treatment of iron deficiency anemias: PANMINERVA MEDICA-EUROPA MEDICA, Vol. 24, No.3. pp.213-220 (July-September 1982).

Folic Acid: Folic Acid is one of the important hematopoietic agents necessary for proper regeneration of the blood-forming elements and their function. Additionally, folic acid increases jejunal glycolytic enzymes and is involved in the desaturation and hydroxylation of long-chain fatty acids in the brain. A deficiency in folic acid results in megaloblastic anemia.

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.

WARNING: Folic acid alone is improper therapy in the treatment of pernicious anemia and other megaloblastic anemias where Vitamin B_{12} is deficient.

PRECAUTIONS: Folic acid in doses above 0.1 mg – 0.4 mg daily may obscure pernicious anemia, in that hematological remission can occur while neurological manifestations remain progressive. The use of this product by immunocompromised patients or treatment of any disorder must be medically supervised by a physician.

DOSAGE AND ADMINISTRATION: Adults (persons over 12 years of age), one (1) capsule daily, between meals, or as prescribed by a physician. Do not exceed recommended dosage. Do not administer to children under the age of 12.

HOW SUPPLIED: Fusion Plus™ are pearl red opaque Vcaps® capsules printed in white with “US” Logo on the cap and “FPlus” on the body. Packed in child resistant cap and light resistant bottle of 30 capsules (52747-0502-30). The listed product number is not a

National Drug Code. Instead, US Pharmaceutical Corporation has assigned this product code formatted according to the standard industry practice to meet the formatting requirements of pharmacy and healthcare insurance computer systems.

CAUTION: Rx only. Rev. 10/2022

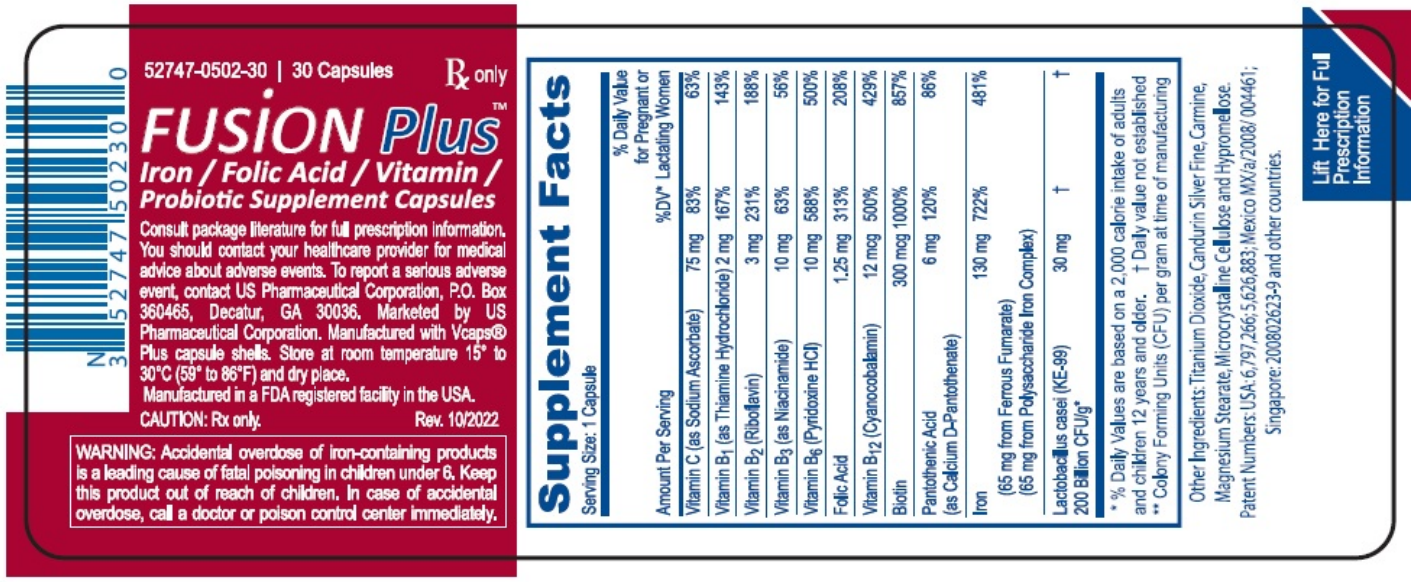
Vcaps® and the Vcaps® Logo are trademarks used under license.

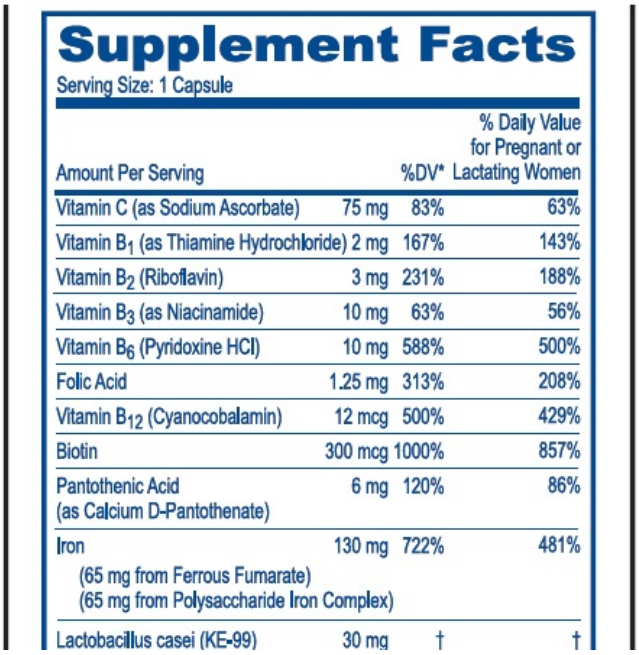
Iron / Folic Acid / Vitamin / Probiotic Supplement Capsules

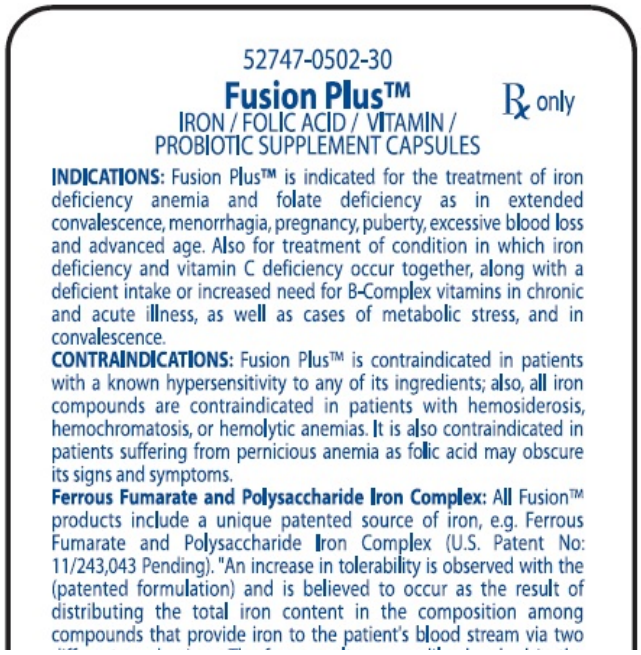
Consult package literature for full prescription information. You should contact your healthcare provider for medical advice about adverse events. To report a serious adverse event, contact US Pharmaceutical Corporation, P.O. Box 360465, Decatur, GA 30036. Marketed by US Pharmaceutical Corporation. Manufactured with Vcaps® Plus capsule shells. Store at room temperature 15° to 30°C (59° to 86°F) and dry place.

Manufactured in a FDA registered facility in the USA.

Packaging







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iron / folic acid / vitamin / probiotic supplement capsule

Product Information			
Product Type	DIETARY SUPPLEMENT	**Item Code (Source)**	NHRIC:52747-502
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ASCORBIC ACID (UNII: PQ6CK8PD0R) (ASCORBIC ACID - UNII:PQ6CK8PD0R)	ASCORBIC ACID	75 mg
THIAMINE (UNII: X66NSO3N35) (THIAMINE ION - UNII:4ABT0J945J)	THIAMINE	2 mg
RIBOFLAVIN (UNII: TLM2976OFR) (RIBOFLAVIN - UNII:TLM2976OFR)	RIBOFLAVIN	3 mg
NIACIN (UNII: 2679MF687A) (NIACIN - UNII:2679MF687A)	NIACIN	10 mg
PYRIDOXINE (UNII: KV2JZ1BI6Z) (PYRIDOXINE - UNII:KV2JZ1BI6Z)	PYRIDOXINE	10 mg
FOLIC ACID (UNII: 935E97BOY8) (FOLIC ACID - UNII:935E97BOY8)	FOLIC ACID	1.25 mg
CYANOCOBALAMIN (UNII: P6YC3EG204) (CYANOCOBALAMIN - UNII:P6YC3EG204)	CYANOCOBALAMIN	0.012 mg
BIOTIN (UNII: 6SO6U10H04) (BIOTIN - UNII:6SO6U10H04)	BIOTIN	0.3 mg
CALCIUM PANTOTHENATE (UNII: 568ET80C3D) (PANTOTHENIC ACID - UNII:19F5HK2737)	PANTOTHENIC ACID	6 mg
IRON (UNII: E1UOL152H7) (IRON - UNII:E1UOL152H7)	IRON	130 mg
LACTICASEIBACILLUS CASEI (UNII: SA940P2U00) (LACTICASEIBACILLUS CASEI -	LACTICASEIBACILLUS	30 mg

UNII:SA940P2U00)	CASEI	30 mg

Inactive Ingredients	
Ingredient Name	**Strength**
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
CARMINIC ACID (UNII: CID8Z8N95N)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NHRIC:52747-502-30	30 in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
dietary supplement		02/01/2013	

Supplement Facts

Serving Size : **Serving per Container :**

Amount Per Serving	% Daily Value
color	
shape	
size (solid drugs) 22 mm	
scoring 1	
imprint	

Labeler - U.S. Pharmaceutical Corporation (079467662)

Revised: 6/2023 U.S. Pharmaceutical Corporation