INTEGRA PLUS - ferrous fumarate and polysacchride iron vitamin mineral complex supplement capsule U.S. Pharmaceutical 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.

see all prescribing information for Integra Plus

DESCRIPTION: Each capsule contains: Ferrous Fumarate (anhydrous)191.1 mg Polysaccharide Iron Complex... 135.9 mg (Equivalent to about 125 mg of elemental iron) Vitamin C (from ProAscorb C‡)....210 mg Folic Acid1 mg Thiamine Mononitrate (B1)...5 mg Riboflavin (B2)....5 mg Niacin (B3) . 20 mg d-Calcium Pantothenate (B5)..7 mg Pyridoxine HCl (B6) ...25 mg Biotin (B7)......300 mcg Cyanocobalamin (B12) .. 10 mcg

CLINICAL PHARMACOLOGY: Integra PlusTM is unique in that it utilizes two (2) different forms of iron, i.e., Ferrous Fumarate and Polysaccharide Iron Complex (as cell-contracted akaganèite), making available a total of 125 mg of elemental iron per capsule as follows:

Ferrous Fumarate (anhydrous) 191.1 mg Polysaccharide iron complex (PIC) 135.9 mg

Subclinical B-Group vitamin deficiencies have greatly increased in recent years, due to changes in dietetic habits, increase in the use of sugar, and the excessive milling of flour and cereals. With thiamine deficiencies in a healthy populous, it is self-evident that the unwell person is particularly prone to thiamine avitaminosis. This is true of the anemic individual with his/her poor appetite and disturbed digestive functions. Folic acid is best known for its role in megaloblastic anemias. Zinc has been recognized in the nutrition of animals and humans, even though the evidence for an uncomplicated zinc deficiency in humans is limited.

Ferrous Fumarate: Provides about 62.5 mg of elemental iron per dose. Ferrous Fumarate is an anhydrous salt of a combination of ferrous iron and fumaric acid, containing 33% of iron per weight. The acute toxicity in experimental animals is low and Ferrous Fumarate is well tolerated clinically. As a ferrous salt, it is more efficiently absorbed in the duodenum. Ferrous Fumarate contrasts very favorably with the availability of the 20% of elemental iron of ferrous sulfate, and the 13% of elemental iron of ferrous gluconate.

Polysaccharide Iron Complex: Provides about 62.5 mg elemental iron, as a cell-contracted akaganèite. It is a product of ferric iron complexed to a low molecular weight polysaccharide. This polysaccharide is produced by the extensive hydrolysis of starch and is a dark brown powder that dissolves in water to form a very dark brown solution, which is virtually odorless and tasteless.

Folic Acid: Folic Acid is one of the important hematopoetic agents necessary for proper regeneration of the blood-forming elements and their function. Folic acid is a precursor of a large family of compounds which serve as coenzymes in carbon transfer reactions. These reactions are required for the synthesis of purine and pyrimidine bases, interconversion of glycine and serine, biosynthesis of methionine methyl groups and degradation of histidine. 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.

All IntegraTM 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 patient's 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: Because Ferrous Fumarate is an organic complex, it contains no free ions, either ferric or ferrous. Polysaccharide Iron Complex is clinically non-toxic. Prior studies in rats demonstrated that Polysaccharide Iron Complex (PIC), administered as a single oral dose to Sprague Dawley rats did not produce evidence of toxicity at a dosage level of 5000 mg Iron/kg: (An Acute Oral Toxicity Study in Rats with Polysaccharide-Iron Complex. T.N.Merriman, M. Aikman and R.E. Rush, Springborn Laboratories. Inc. Spencerville, Ohio Study No. 3340.1 March - April 1994). Other clinical studies had demonstrated that Polysaccharide Iron gives a good hematopoietic response with an almost complete absence of the side effects usually associated with oral iron therapy. Picinni and Ricciotti suggested in 1982, that "the therapeutic effectiveness of Polysaccharide Iron Complex when compared with iron fumarate in the treatment of iron deficiency anemia, appears to be as active as the iron fumarate and as well tolerated. however, it exerted a greater influence on the level of hemoglobin and on the number of red cells..." and that, "it has been exceptionally well tolerated by all patients" (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).

As mentioned above, the patented source of iron used in Integra PlusTM (Ferrous Fumarate and Polysaccharide Iron Complex) provides a high level of elemental iron with a low incidence of gastric distress.

CONCLUSION: Based on the results of this study, the oral combination of Ferrous Fumarate and Polysaccharide Iron Complex was better tolerated and safer than the oral administration of Ferrous Fumarate alone. The conclusion of this research stated, that the addition of PIC to Ferrous Fumarate surprisingly allows the same concentration of Ferrous Fumarate to be better tolerated than the Ferrous Fumarate alone.

CONTRAINDICATIONS: Integra PlusTM 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. Pernicious anemia is a contraindication, as folic acid may obscure its signs and symptoms.

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 B12 is deficient.

PRECAUTIONS: General: 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. Pediatric Use: Safety and effectiveness of this

product have not been established in pediatric patients. Geriatric Use: No clinical studies have been performed in patients age 65 and over to determine whether older persons respond differently from younger persons. Dosage should always begin at the low end of the dosage scale and should consider that elderly persons may have decreased hepatic, renal, or cardiac function and or concomitant diseases.

Adverse Reactions: Folic Acid: Allergic sensitizations have been reported following both oral and parenteral administration of folic acid. Ferrous Fumarate: 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 Integra PlusTM after meals may control occasional G.I. disturbances. Integra PlusTM is best absorbed when taken at bedtime.

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. Other symptoms include pallor and cyanosis, melena, shock, drowsiness and coma. The estimated overdose of orally ingested iron is 300-mg/kg body weight. When overdoses are ingested by children, severe reactions, including fatalities, have resulted. Integra PlusTM should be stored beyond the reach of children to prevent against accidental iron poisoning. Keep this and all other drugs out of the reach of children.

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

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.

Pediatric Use: Safety and effectiveness of this product have not been established in pediatric patients.

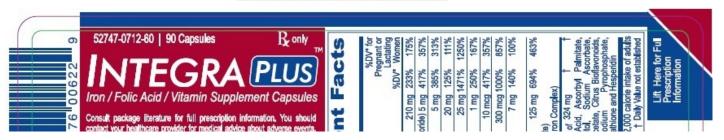
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HOW SUPPLIED: Integra PlusTM are Vcaps[®] capsules with maroon caps printed in white with "Integra Plus" and cream body printed in red with "US" logo. Packed in child resistant caps and light resistant bottles of 90 capsules (52747-0712-60) and 30 capsules (52747-0712-30). The listed product numbers are not National Drug Codes. Instead, US Pharmaceutical Corporation has assigned these product codes formatted according to standard industry practice to meet the formatting requirements of pharmacy and healthcare insurance computer systems.

CAUTION: Rx only.

Packaging

PACKAGE 90 CAPSULES





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 (58° to 86°F) and dry place. Manufactured in a FDA registered facility in the USA.

CAUTION: Rx only.

Vcaps® and the Vcaps® Logo are trademarks used under license. Patent Numbers: USA: 5,626,883; Mexico MX(a/2008/004461; Singapore: 200802623-9 and other countries.

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.

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Amount Per Serving Vitamin C (from ProAscorb C Vitamin B1 (as Thiamine Hyd Vitamin B2 (Riboflavin) Vitamin B3 (as Niacinamide) Vitamin B6 (as Pyridoxine HC Folic Acid Vitamin B12 (Cyanocobalami Biotin Pantolihenic Acid	(as Calcium d-Pantothenate) Iron
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Other Ingredients: Hypromellose, Magnesium Stearate, Titanium Dioxide, FDA/E172 Yellow Iron Oxide, FD&C Red #40, and FD&C Blue #1.

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Supplement Facts

Serving Size: 1 Capsule			
Amount Per Serving		Pre	%DV* for egnant or Lactating Women
Vitamin C (from ProAscorb C**)	210 mg	233%	175%
Vitamin B ₁ (as Thiamine Hydrochk	oride) 5 mg	417%	357%
Vitamin B ₂ (Riboflavin)	5 mg	385%	313%
Vitamin B ₃ (as Niacinamide)	20 mg	125%	111%
Vitamin B ₆ (as Pyridoxine HCI)	25 mg	1471%	1250%
Folic Acid	1 mg	250%	167%
Vitamin B ₁₂ (Cyanocobalamin)	10 mcg	417%	357%
Biotin	300 mcg	1000%	857%
Pantothenic Acid (as Calcium d-Pantothenate)	7 mg	140%	100%
Iron	125 mg	694%	463%

(62.5mg from Ferrous Fumarate) (62,5mg from Polysaccharide Iron Complex)

**ProAscorb C is a proprietary blend of 324 mg Calcium Ascorbate, Ascorbic Acid, Ascorbyl Palmitate, Niacinamide Ascorbate, Xylitol, Sodium Ascorbate, Magnesium Ascorbate, Lysine Acetate, Citrus Bioflavonoids, Potassium Ascorbate, Sodium Pyrophosphate, Potassium Ascorbate, Sodium Pyrophosphate, D-Ribofuranose, L-Cysteine, Glutathione and Hesperidin

* % Daily Values are based on a 2,000 calorie intake of adults and children 12 years and older. † Daily Value not established

52747-0712-60 Integra Plus ™

R only

IRON / FOLIC ACID / VITAMIN SUPPLEMENT CAPSULES

INDICATIONS: Integra 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, as well as cases of metabolic stress, and in convalescence.

CONTRAINDICATIONS: Integra 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: Al Integra™ 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 patient's 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

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WARNING: Folic acid alone is improper therapy in the treatment of pernicious anemia and other megaloblastic anemias where Vitamin B₁₂ is

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.

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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: Integra Plus™ are Vcaps® capsules with maroon caps printed in white with "Integra Plus" and cream body printed in red with "US" logo. Packed in child resistant caps and light resistant bottles of 90 capsules (52747-0712-60) and 30 capsules (52747-0712-30). The listed product numbers are not National Drug Codes. Instead, US Pharmaceutical Corporation has assigned these product codes formatted according to standard industry practice to meet the formatting requirements of pharmacy and healthcare insurance computer systems.

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PACKAGE 30 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, Po. Box 360466, Decatur, GA 30036. Marketed by US Pharmaceutical Corporation. Manufactured with Vosps® 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.

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Vcaps® and the Vcaps® Logo are trademarks used under license. Patent Numbers: USA: 5,626,883; Mexico MX/a/2008/004461; Singapore: 200802623-9 and other countries.

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Facts

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Serving Size: 1 Capsule

m B only

125% 250% 417% 1000% 20 mg 25 mg 1 1 mg 10 mcg 300 mcg 1 7 mg 5 mg

(as Pyridoxine HCI)

Vitamin B6 (Folic Acid

Vitamin B₃ (as Niacinamide)

Vitamin B2 (Riboflavin)

Vitamin B₁₂ (Cyanocobalamin)

ntothenic Acid Calcium d-Pantothenate)

Par

Vitamin B₁ (as Thiamine Hydrochloride) 5 mg

Vitamin C (from ProAscorb C**)

Amount Per Serving

694% mg 25

"ProAscorb C is a proprietary blend of 324 in Calcium Ascorbate, Ascorbic Acid, A Viacinamide Ascorbate, Xylitol, Si from Ferrous Furnarate) from Polysaccharide Iro 5mg 5mg 85.55 (8)

based on a 2,000 calone intake of adults and older. † Daily Value not established Lysine

Sodium Ascorbate Citrus Bioflavonoids

Sodium Pyrophosph Glutathione and Hesperidin

Lift Here for Full Prescription Information

Inactive Ingredients: Hypromellose, Magnesium Stearate, Titanium Dioxide, FDA/E172 Yellow Iron Oxide, FD&C Red #40, and FD&C Blue #1.

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52747-0712-30

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INTEGRA PLUS

ferrous fumarate and polysacchride iron vitamin mineral complex supplement capsule

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:52747-712
Route of Administration	ORAL		

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
FERROUS FUMARATE (UNII: R5L488RY0Q) (FERROUS CATION - UNII:GW89581OWR)	FERROUS CATION	62.5 mg		
FERROUS ASPARTO GLYCINATE (UNII: H7426RGB3L) (FERROUS CATION - UNII:GW895810WR)	FERROUS CATION	62.5 mg		
FOLIC ACID (UNII: 935E97BOY8) (FOLIC ACID - UNII:935E97BOY8)	FOLIC ACID	1 mg		
ASCORBIC ACID (UNII: PQ6CK8PD0R) (ASCORBIC ACID - UNII:PQ6CK8PD0R)	ASCORBIC ACID	210 mg		
NIACIN (UNII: 2679MF687A) (NIACIN - UNII:2679MF687A)	NIACIN	20 mg		
THIAMINE MONONITRATE (UNII: 8K0I04919X) (THIAMINE ION - UNII:4ABT0J945J)	THIAMINE	5 mg		
RIBOFLAVIN (UNII: TLM29760FR) (RIBOFLAVIN - UNII:TLM29760FR)	RIBOFLAVIN	5 mg		
CALCIUM PANTOTHENATE (UNII: 568ET80C3D) (PANTOTHENIC ACID - UNII:19F5HK2737)	PANTOTHENIC ACID	5 mg		
PYRIDOXINE HYDROCHLORIDE (UNII: 68Y4CF58BV) (PYRIDOXINE - UNII: KV2JZ 1BI6Z)	PYRIDOXINE HYDROCHLORIDE	25 mg		
BIOTIN (UNII: 6SO6U10H04) (BIOTIN - UNII:6SO6U10H04)	BIOTIN	300 ug		

Inactive Ingredients	
Ingredient Name	Strength
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
FERRIC OXIDE YELLOW (UNII: EX43802MRT)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Product Characteristics				
Color	red (Maroon body and Yellow capsule)	Score	no score	
Shape	CAPSULE	Size	22mm	
Flavor		Imprint Code	Integra;Plus;US	
Contains				

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52747- 712-60	90 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/27/2009	
2	NDC:52747- 712-30	30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/27/2009	
3	NDC:52747- 712-10	40 in 1 BOX	04/27/2009	
3	NDC:52747- 712-04	4 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information				
Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
	04/27/2009			
	Application Number or Monograph	Application Number or Monograph Marketing Start Citation Date		

Labeler - U.S. Pharmaceutical Corporation (079467662)

Registrant - U.S. Pharmaceutical Corporation (079467662)

Revised: 4/2024 U.S. Pharmaceutical Corporation