MULTIVITAMINS + IRON

REGERON™ - E PLUS

Capsule

VITAMINS AND MINERALS

FORMULATION:

Each capsule contains:

Alpha Tocopherol (Vit. E)	200 mg
Thiamine HCl (Vit. B1)	10 mg
Riboflavin (Vit. B2)	5 mg
Pyridoxine HCl (Vit. B6)	10 mg
Cyanocobalamin (Vit. B12)	10 mcg
Ascorbic Acid (Vit. C)	50 mg
Calcium Pantothenate	5 mg
Nicotinamide	10 mg
Ferrous Sulfate-anhydrous	54.4 mg
(Equivalent to 20 mg Elemental Iron)	
Buclizine HCl	25 mg

PRODUCT DESCRIPTION:

REGERON™ - E PLUS Capsule is a light orange to yellow-orange granules in a brownish violet/brownish violet capsule #0

This particular formulation of MULTIVITAMINS + IRON (**REGERON™ - E PLUS**) Capsule is a comprehensive nutritional supplement incorporating substantial amounts of various essential vitamins fortified with elemental iron and an appetite stimulant.

PROPERTIES AND ACTIONS:

The wide-ranging formulation of this special preparation of MULTIVITAMINS + IRON (**REGERON™ - E PLUS**) Capsule guarantees broad, effective and reliable nutritional support. Specifically, it enhances food intake, induces weight gain, supplements inadequate vitamin and iron intake, and improves metabolism of nutrients. The sweeping consequent benefits to patients include improved bodily functions, a general sense of well-being, and enhanced performance capacities during increased physical and mental activities.

INDICATIONS:

For stimulation of appetite specially in conditions associated with underweight cases, malnutrition, malaise, easy fatigability and general body weakness.

As nutritional support in chronic illness, decreased resistance to infection, old age and debility, convalescence after illness or surgery, and in restrictive or deficient diet.

DOSAGE AND ADMINISTRATION:

One capsule daily or as prescribed by the physician.

CONTRAINDICATIONS:

There are no known contraindications.

PRECAUTIONS:

Owing to its possible sedative effect, patients may slow their reactions when driving vehicles, and operating machines.

Should not be taken simultaneously with alcohol or other CNS depressants.
The safety of buclizine in pregnancy has not been determined.

ADVERSE DRUG REACTION:

Mild drowsiness may occur at the beginning of treatment which gradually disappears after 2-3 days of continuous therapy.

Occasional gastrointestinal discomfort such as nausea and diarrhea may occur with intake of iron.

DRUG INTERACTIONS:

Concomitant administration of tetracycline and some antacids may decrease the amount of iron absorbed and similarly, iron may retard the absorption of tetracyclines. These products should not be taken within 2 hrs. of each other. Food reduces the absorption of iron but will lessen gastric irritation.

OVERDOSAGE AND TREATMENT:

Symptoms: Drowsiness, nausea, hypertension, dizziness, excitatory states (in children), respiratory depression, convulsion (particularly in children) and coma.

Treatment: Gastric lavage followed by administration of activated charcoal. If necessary, benzodiazepines and symptomatic treatment including cardiovascular and respiratory monitoring for excitatory states or convulsions must be given.

CAUTION:

Foods, Drugs, Devices and Cosmetics Act prohibits dispensing without prescription.

AVAILABILITY:

Amber blister pack of 10's (Box of 50's)

STORE AT TEMPERATURES NOT EXCEEDING 30°C.

For suspected adverse drug reaction, report to the FDA:
www.fda.gov.ph

DR-XY11925

SVMore

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