Name of medicine

Hydroxocobalamin (vitamin B<sub>12</sub>) as chloride.
Molecular formula: C<sub>62</sub>H<sub>90</sub>ClCoN<sub>13</sub>O<sub>15</sub>P.
Molecular weight is 1,383.

Description

Hydroxocobalamin is a dark red, odourless crystalline powder or crystals. It is soluble in water and alcohol, sparingly soluble in methyl alcohol and practically insoluble in acetone, chloroform and ether. HYDROXO-B12 also contains sodium chloride 9.0 mg and hydrochloric acid for pH adjustment and water for injections. The pH of the solution is approximately 4.6.

Pharmacology

Several chemically related forms of vitamin B<sub>12</sub>, differing in slight modification of a side chain attached to the cobalamin nucleus have been isolated. Two such variants of vitamin B<sub>12</sub> are cyanocobalamin and hydroxocobalamin. Vitamin B<sub>12</sub> is essential for normal growth, haemopoiesis, production of all epithelial cells and maintenance of myelin throughout the nervous system. Whenever nucleic acid synthesis occurs
and therefore whenever cell reproduction occurs, vitamin B₁₂ is required. The amounts of vitamin B₁₂ needed to maintain normal blood forming functions are small and low doses are sufficient to correct the usual symptoms of vitamin B₁₂ deficiency. Vitamin B₁₂ acts as an enzyme or coenzyme in a number of metabolic processes and is transformed in the body to at least two compounds which possess enzymatic properties.

i) Coenzyme B₁₂ is required for conversion of propionate to succinate, thus involving vitamin B₁₂ in both fat and carbohydrate metabolism.

ii) Methylcobalamin acts in a transmethylation process converting homocysteine to methionine, thus involving vitamin B₁₂ in fat and protein metabolism. In some cases of vitamin B deficiency, severe neurological symptoms develop, as vitamin B₁₂ is necessary for the formation of protein structures required for the integrity of the nerve cell and myelin sheath.

**Pharmacokinetics**

Hydroxocobalamin produces higher and more prolonged serum levels of vitamin B₁₂ than cyanocobalamin when given by intramuscular injection in the same dosage. Hydroxocobalamin disperses more slowly from the site of injection than cyanocobalamin, is more strongly bound to plasma proteins and accumulates in the liver to a greater extent. Hydroxocobalamin is excreted in the bile and urine, but more slowly than cyanocobalamin. Hydroxocobalamin combines with cyanide and thus acts as a cyanide antagonist *in vivo* resulting in the formation of cyanocobalamin.

**Indications**

Prophylaxis and treatment of pernicious (Addisonian) anaemia and other macrocytic anaemias associated with vitamin B₁₂ deficiency. Treatment of optic neuropathies such as tobacco amblyopia and Leber's optic atrophy.

**Contraindications**

Known sensitivity to hydroxocobalamin or any other ingredient in HYDROXO-B12. Known sensitivity to cobalt. Hydroxocobalamin should not be used for the treatment of megaloblastic anaemia of pregnancy (see Precautions, Use in pregnancy).

**Precautions**

Do not use intravenously. A sensitivity history should be obtained from the patient prior to administration of vitamin B₁₂. An intradermal test dose is recommended before vitamin B₁₂ is administered to patients who may be sensitive to cobalamins.

Hypokalaemia and cardiac arrest have been reported when megaloblastic anaemia is treated intensively.
Serum potassium is to be carefully monitored during the initial phase of treatment in pernicious anaemia.

Diagnosis of vitamin B\textsubscript{12} deficiency should be confirmed by laboratory investigation before institution of hydroxocobalamin (vitamin B\textsubscript{12}) therapy.

Do not use hydroxocobalamin until diagnosis is fully established, as it may mask symptoms of subacute degeneration of the spinal cord, or of the true diagnosis of pernicious anaemia. Folic acid may potentiate the neurological complications of vitamin B\textsubscript{12} deficiency, so it should not be administered to patients with pernicious anaemia (see Interactions). Regular blood tests to determine vitamin B\textsubscript{12} levels are advisable during treatment. Administration of hydroxocobalamin doses in excess of 10 microgram daily may improve folate deficient megaloblastic anaemia and obscure the true diagnosis.

The therapeutic response to hydroxocobalamin may be impaired by concurrent infection, uraemia, folic acid or iron deficiency, or by drugs with bone marrow suppressing effects, such as chloramphenicol (see Interactions).

Treatment with hydroxocobalamin may unmask polycythaemia vera, because vitamin B\textsubscript{12} deficiency may suppress the symptoms of this condition.

**Use in pregnancy**
Problems in humans have not been documented with intake of normal daily amounts. Vitamin B\textsubscript{12} crosses the placental barrier. There are no studies establishing the safety of this drug during pregnancy. It is not recommended for pregnancy unless the expected benefits outweigh any potential risk to the infant. Megaloblastic anaemia occurring during pregnancy is usually due to folic acid deficiency rather than vitamin B\textsubscript{12} deficiency.

Hydroxocobalamin should not be used for the treatment of megaloblastic anaemia of pregnancy caused by folic acid deficiency.

**Use in lactation**
Hydroxocobalamin is distributed into breast milk. Therefore it is not recommended for breastfeeding mothers unless the expected benefits to the mother outweigh any potential risk to the infant.

**Interactions**
Concurrent administration of chloramphenicol and hydroxocobalamin may impair the therapeutic response to hydroxocobalamin in vitamin B\textsubscript{12} deficient patients. The haematological response should be carefully monitored in patients receiving both these drugs. Serum concentrations of hydroxocobalamin may be lowered by oral contraceptives. Vitamin B\textsubscript{12} concentrations in the blood may be reduced following administration of large and continuous doses of folic acid. Folic acid administration may impair the therapeutic response to hydroxocobalamin.

**Laboratory tests**
Most antibiotics, methotrexate and pyrimethamine invalidate folic acid and vitamin B\textsubscript{12} microbiological blood assays.
**Adverse effects**

Sensitisation to hydroxocobalamin is rare, but may manifest itself as itching exanthema and rarely, anaphylaxis. Antibodies to hydroxocobalamin transcobalamin II complex may develop during hydroxocobalamin therapy.

Other reported adverse effects include diarrhoea, nausea, vomiting, headache, dizziness, peripheral vascular thrombosis, chest pain/discomfort, cardiac arrest, injection site reactions, sensation of heat and cold, malaise, urticaria or a feeling of swelling of the whole body, eczematous skin lesions, acne and folliculitis.

Pulmonary oedema and congestive heart failure have been reported during early vitamin B\(_{12}\) treatment, possibly as a result of an increase in blood volume induced by the drug.

Polycythaemia vera may occur (see Precautions).

Arrhythmias secondary to hypokalaemia have appeared at the beginning of parenteral treatment with hydroxocobalamin.

**Dosage and administration**

This product contains no antimicrobial agent. It is for single use in one patient only. Discard any residue. Hydroxo-B12 Injection is to be administered intramuscularly.

The following dosage schemes are suitable for adults and children:

**Addisonian pernicious anaemia and other macrocytic anaemias without neurological involvement.**

*Initially.* 250 to 1,000 micrograms intramuscularly on alternate days for one to two weeks, then 250 micrograms weekly until the blood count is normal.

*Maintenance.* 1,000 micrograms every two or three months.

**Addisonian pernicious anaemia and other macrocytic anaemias with neurological involvement.**

*Initially.* 1,000 micrograms on alternate days for one to two weeks.

*Maintenance.* 1,000 micrograms every two months.

**Prophylaxis of macrocytic anaemia associated with vitamin B12 deficiency resulting from gastrectomy, some malabsorption syndromes and nutritional deficiencies.**

1,000 micrograms every two or three months.
Tobacco amblyopia and Leber's optic atrophy.

*Initially.* 1,000 microgram daily by intramuscular injection for two weeks then twice weekly for four weeks.  
*Maintenance.* 1000 microgram monthly.

**Presentation and storage conditions**

Solution for injection, 1mg/mL (1000 microgram/mL) (dark red, clear): 3’s (ampoules).  
Store below 25°C  
Protect from light

**Poison Schedule**

Unscheduled

**Sponsor**

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