

STANOZOLOL DEPOT

STANOZOLOL FOR INJECTION USP 50mg Water Base q.s. Presented as 10 x 1ml ampoules per box (50mg/ml)

DESCRIPTION

Stanozolol Injection is indicated prophylactically to decrease the frequency and severity of attacks of angioedema. It is also indicated breast cancer in post-menopausal women, for anaemias, osteoporosis, and catabolic disorders.

Stanozolol Injection is a white suspension for Intramuscular use. Each ml contains Stanozolol 50 mg. Stanozolol is Odorless, crystalline powder with empirical formula C21H32N2O.

PHARMACODYNAMICS

Anabolic steroids are synthetic derivatives of Testosterone. Stanozolol has been found to increase low-density lipoproteins and decrease high-density lipoproteins. These changes are not associated with any increase in total cholesterol or triglyceride levels and revert to normal on discontinuation of treatment. Hereditary angioedema (HAE) is an autosomal dominant disorder caused by a deficient or non-functional C1 esterase inhibitor (C1 INH) and clinically characterized by episodes of swelling of the face, extremities, genitalia, bowel wall, and upper respiratory tract.

PHARMACOKINETICS

Reverses catabolic processes and negative nitrogen balance by promoting protein anabolism and stimulating appetite if there is concurrently a proper intake of calories and proteins.

INDICATIONS AND USAGE

Hereditary Angioedema: Stanozolol is indicated prophylactically to decrease the frequency and severity of attacks of angioedema. It is also indicated breast cancer in postmenopausal women, for anaemias, osteoporosis, and catabolic disorders.

DOSAGE AND ADMINISTRATION

The use of anabolic steroids may be associated with serious adverse reactions, many of which are dose related. Therefore patients should be placed on the lowest possible effective dose. Stanozolol has been given by intramuscular injection in doses of 50 mg to 100mg every 1 or 3 times per week. The prophylactic dose of Stanozolol to be used prior to dental extraction or other traumatic or stressful situations has not been established and may be substantially larger.

CONTRAINDICATIONS

The use of Stanozolol is contraindicated in the following: Male patients with carcinoma of the breast or with known or suspected carcinoma of the prostate.

Carcinoma of the breast in females with hypercalcemia; androgenic anabolic steroids may stimulate osteolytic resorption of bone.

Nephrosis or the nephrotic phase of nephritis. Stanozolol can cause fetal harm when administered to a pregnant woman.

Genitourinary System:

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In men, Prepubertal: Phallic enlargement and increased frequency of erections. Postpubertal: Inhibition of testicular functions, testicular atrophy and oligospermia, impotence, chronic priapism, epididymitis and bladder irritability.

In women: Clitoral enlargement, menstrual irregularities.
In both sexes: increased or decreased libido. CNS: Habituation, excitation, insomnia, and depression.

Gastrointestinal: Nausea, vomiting, diarrhea.

Hematologic: Bleeding in patients on concomitant anticoagulant therapy.

Breast: Gynecomastia. Larynx: Deepening of the voice in women. Hair: Hirsutism and male pattern baldness in women. Skin: Acne (especially in women and prepubertal boys). Skeletal: Premature closure of epiphyses in children. Fluid and Electrolytes: Edema, retention of serum electrolytes (Sodium, chloride, potassium, phosphate, and calcium).

OVERDOS

An overdose of this medication is unlikely to threaten health. Symptoms of Stanozolol overdose are not known.

PRESENTATION

50 mg/ml, 1 x 10 ml ampoules

STORAGE

Store in a cool dry place (30 °C ± 2 °C). Protect from light. Warming and rotating the ampoule between the palms of the hands will redissolve any crystals that may have been formed during storage at low temperatures.

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