ketoconazole

therapy

Vaginal candidosis:	In all dermatological and in systemic fungal infections:
2 tablets once daily (with food) for 5 days	1 tablet daily (with food) until complete sympto- matic and mycological cure is obtained

Not all indications are as yet approved in all countries.

PRESCRIBING INFORMATION

Presentation: white flat half scored uncoated tablets marked "Janssen" on one side and k/200 on the reverse. Each tablet contains 200 mg ketoconazole. Uses: Nizoral is an orally active antimycotic for the treatment in adults, of vaginal candidosis, superficial and systemic mycoses including dermatophyte and yeast infections of the skin, hair and nails, yeast infections of the mouth and G.I. tract. Also maintenance treatment of systemic mycoses and chronic mucocutaneous candidosis and prophylaxis in chronic mucocutaneous candidosis and prophylaxis in at risk: patients. In children: systemic mycoses and severe local infections where previous topical treatment has failed.
Side-effects, precautions, contra-indications: contra-indicated in pregnancy. For maximal absorption. Nizoral should be taken with meals. The use of agents which reduce asstric acidity (articholiperaic drugs antacids. He blockers). gastric acidity (anti-cholinergic drugs, antacids, H₂ blockers) should be avoided and, if indicated, such drugs should be taken not less than two hours after **Nizoral**. Nausea, skin taken not less than two hours after **Nizoral**. Nausea, skin rash, headache and pruritus may occasionally be observed. Alterations in liver function tests have occurred in patients on ketoconazole, these changes may be transient. Cases of hepatitis have been reported with an incidence of about 1 per 10,000 patients. Some of these may represent an idiosyncratic adverse reaction to the drug. This should be borne in mind in patients on long-term therapy. If a patient develops jaundice or any symptoms suggestive of hepatitis, treatment with ketoconazole should be stopped.



Janssen Pharmaceutica B-2340 Beerse, Belgium

STATISTICS by T. D. V. SWINSCOW T SOUARE ONE

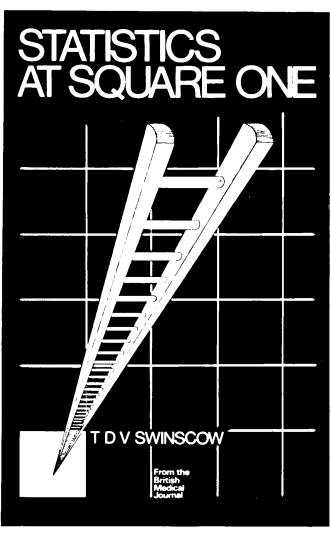
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O'Donnell, Barry, British Medical Journal, 1977, 1, 451.

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Prescribing Information

Uses Management of Diabetes mellitus.

Dosage and administration Dosage to be determined by the physician. Site of injection to be changed according to suitable routine. Avoid unintentional intravascular

Neusulin, Soluble Insulin: Administered s.c., i.m. or i.v. S.c., onset of action within 30-60 minutes, duration 6-8 hours. I.m., onset is faster and duration is shorter. I.v. administration has fastest onset and shortest duration, usually reserved for investigational use or diabetic ketoacidosis.

Neuphane, Neulente: Administered s.c. or i.m. Not to be given i.v. S.c., onset of action within 2 hours, duration (Neuphane) 20-24 hours, duration (Neulente) 24-28 hours. I.m., onset is faster and duration shorter. Mix well by gently inverting the vial several times before use.

Mixing: Neusulin may be mixed in the syringe, on medical advice, with Neuphane or Neulente if required, provided the mixture is injected immediately. However, it is preferable to avoid mixing insulins of different pH.

See data sheet for procedure.

Contra-indications Hypoglycaemia.

Precautions Dosage requirement may alter with variation of lifestyle, infection, pregnancy and with change in

species, type or purity of insulin. Hypo- and hyperglycaemia may be enhanced by drugs which interact with insulin. Beta-blockers may affect insulin requirement and mask hypoglycaemia. MAO

inhibitors may potentiate insulin.

Side-effects Hypoglycaemia. Possible altered visual refraction. Transient local reactions at the site of injection. **Storage** Store at 2-8°C. *Do not freeze*. Avoid direct sunlight. Presentation Neusulin, Neuphane and Neulente are available in strengths of 40 and 80 units per ml in vials of 10 ml.

Basic NHS costs

Neusulin 40 units/ml PL3/0137 £2.31 80 units/ml PL3/0138 £4.14

40 units/ml PL3/0141 £2.28 80 units/ml PL3/0142 £3.90

Neuphane

40 units/ml PL3/0139 £2.31 80 units/ml PL3/0140 £4.47

Further information is available on request.

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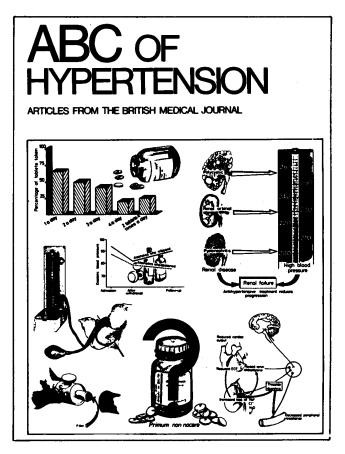
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Voltarol[®] for estecarthritis

Presentation Tablets of 25 and 50 mg diclofenac sodium; suppositaries at 100 mg diclofenac sodium. Indications Rheumatoid arthriffs, asteoarthriffs, low back pain and other ocute muscula-stational disorders, ankylosing spondyliffs, acute gout. Dosage Tablets: 75-150 mg daily in two or three divided doses. Suppositaries: one daily, usually at night. Interiore severe cases, combined therapy with tablets is recommended (daily dose should not exceed 150 mg). Side-effects Transient epigastric point, natifica and diarrhoea, headlache and slight dizziness have been reported, as occasionally have skin rash, peripheral oxidema and abnormalities of serum transaminases and (very rarely) peptic ulcer and haematemesis or melaena. Local reactions to suppositories include itching, burning and increased traquency, of bipwel movement. Precautions Do not prescribe during pregnancy unless there are compelling reasons. Patients with a history of peptic ulcer, haematemesis or melaena, or with severe hepatic or renal insufficiency, should be kept under close surveillance. Voltarol has been reported to depress solicylate levels and vice versa: the clinical relevance of this is not yet clear. Use suppositaries only with caution in patients with pointing in rivitable ano-rectal conditions. Contro-indications Peptic ulceration; patients in whom attacks of asthma, unicaria or acute rhinitis are precipitated by ospirin or other NSAIs. Packs Toblets. 25 mg (PL0001/0083) in packs of 100, basic NHS seems cribing information is dvailable from Geigy Pharmaceuticals, Horsham, West Sussex.