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Prescribing information

resentation: white, flat, half scored uncoated tablets marked "Janssen" on one side and K/200 on the reverse. Each tablet contains

200 mg ketoconazok

Uses: **illustratic** is an orally active antiniy cotic for the treatment in adults of vaginal candidosis, superficial and systemic mycoses including dermatophyte and yeast infections of the mouth and G.I.-tract. Also maintenance treatment of systemic mycoses and chronic micocutaneous 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-inflications: contra-indicated in pregnancy. For maximal absorption libroral should be taken will meals. The use of agents winch redule gestra exidity functionliency derives, antacids. Hy, biotecers should be avoided and, if indicates such drugs should be taken not less than two hours after libroral shuses, skin rash, beausite and pruritus may occasionally be observed Alterations include for each state of the patients have been shared to a state of the patients and a state of the patients and the patients are stated experted 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-time therapy. If a patient develops jaundice or any symptom suggestive or hepatitis treatment with ketoconazoie shouldbe stopped sillo asymptomatic increases of liver enzyme levels, on the other hand, do not necess tate discontinuation of the treatment. Ketoconazoie, when given together with cyclosporin A results in increased blood levels of cyclosporin A are monitored if the two drugs are given concomitantly, ketoconazoi 200 mg once daily produces a transient decrease in plasma levels of testosterone, During long term therapy at this dose, testosteron levels are usually not significantly different from controls in rare instances, gynaecomastic has been reported.



#### Heceta lighthouse, Oregon, USA

#### Prescribing Information

Presentation Vials containing 500mg, 1g or 2g of cefotaxime sodium. Indications Infections before identification of the organism. Infections caused by bacteria of established sensitivity, including chest infections, septicaemia, urinary tract infections, soft tissue infections, obstetric and gynaecological infections, bone and joint infections, meningitis, gonorrhoea. Dosage Claforan is administered i.m. or i.v. Adults: Usually 2-8g daily (see full prescribing information). For infections caused by sensitive Pseudomonas spp., doses of more than 6g daily are usually required. Children: 100-150mg/kg/day may be given in very severe infections. Neonates: 50mg/kg body weight daily in 2 to 4 equally divided doses. In cases of severe infection, divided daily doses of 150-200mg/kg have been given. Dosage in renal impairment Reduced dosage is only required in severe enal failure (GFR<5ml/min = serum creatinine approx. 751 µmol/l) when, after an initial loading dose of 1g, the daily dose is halved without change in frequency of dosing. Contra-Indications Known allergy to cephalosporins. Precautions Cephalosporin antibiotics may usually be given safely to patients who

are hypersensitive to penicillins, although cross reactions have been reported. Special care is indicated in patients who have had an anaphylactic response to penicillin. Patients with severe renal dysfunction — see previous. Cephalosporin antibiotics at high dosage should be given with caution to patients receiving aminoglycoside antibiotics or potent diuretics such as frusemide. At recommended doses, enhancement of nephrotoxicity is unlikely with Claforan. A false-positive reaction to glucose may occur with reducing substances. Claforan should not be mixed in the syringe with aminoglycoside antibiotics. The safety of Claforan in human pregnancy has not been established. Side effects Adverse reactions are rare and generally mild and transient, but include diarrhoea (pseudomembraneous colitis has been rarely reported), candidiasis, rashes, fever, eosinophilia, leukopenia, transient rises in liver transaminase and alkaline phosphatase, transient pain at the site of injection and phlebitis. Product licence holder and number Roussel Laboratories Ltd., Broadwater Park, North Orbital Road, Uxbridge, Middlesex UB9 5HP. 0109/0074. Package quantities and basic N.H.S. price Vials of 500mg, 1g and 2g in packs of 10. One gram vial £4.95. Date of preparation November 1984.



Publication date 21 May 1985

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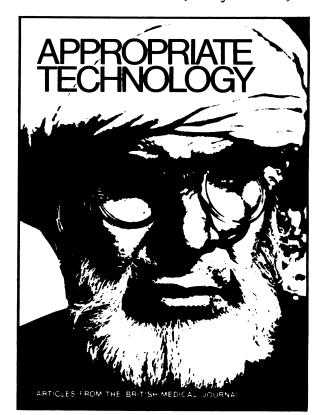
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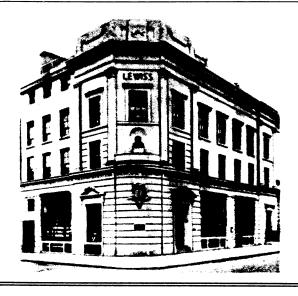


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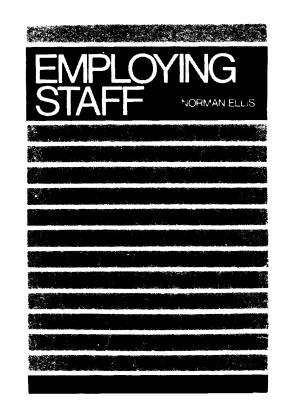
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Often moving, occasionally horrific, sometimes hilarious, these highly personal memories reflect the many aspects of war from a medical viewpoint.

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