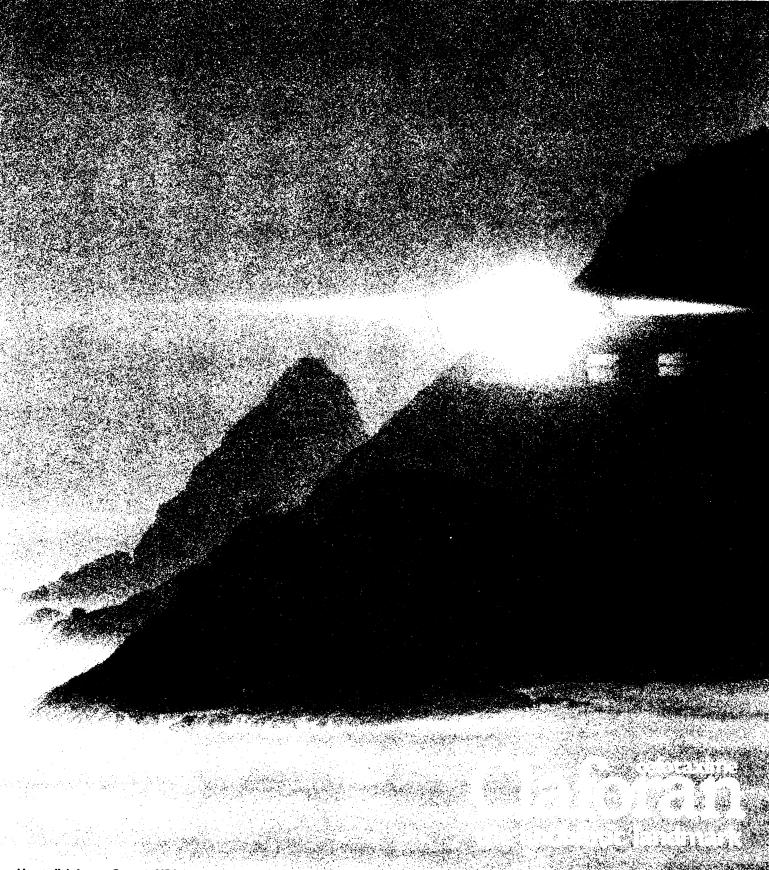


islike of topical application the patie leads to in ate therapy

Presentation: white, flat, half scored uncoated tablets marked "Janssen" on one side and K/200 on the reverse. Each tablet contains 200 mg (encoronazole. Uses: National is an orally active antimycotic for the treatment in adults of vaginal candidosis, superficial and systemic mycoses including dematophyte and yeast infections of the skin, hair and nalis, yeast infections of the mouth and G1-tract. Also maintenance treatment of systemic mycoses and chronic mucocutaneous candidosis and prophylaxis in "at risk", patents: In children: systemic mycoses and severe local infections where previous topical treatment has failed. Side-effects, precaustions, contral-indicatedions: contral-indicated in pregnancy. For maximal absorption Nationard should be taken with meals. The use of agents which reduce gastric acidity (anti-cholinergic drugs, antacids, Hy-blockers) should be avoided and. If indicated, such drugs should be taken not less than two hours after Nationardi. Nausea, skin rash, headache and pruntus 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 indiopyncratic 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, Mild asymptomatic increases of liver enzyme levels, on the other hand, do not necessitate discontinuation of the treatment, Ketoconazole, when given together with cyclosponin A results in increased blood levels of cyclosponin A, its important that blood levels of cyclosponin A are monitored in the wordings are given concomitantly. Ketoconazole. During long term therapy at this dose, testosterone levels are usually not significantly different from controls. In rare instances, gynaecomastia 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, obsettric and gynaecological infections, bone and joint infections, meningitis, gonorrhoea. Dosage Claforan is administered i.m. or i.v. Adults: Usually 2-6g 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 in 2 to 4 divided doses. Up to 200mg/kg/day may be given in very severe infection, divided daily doses of 500-200mg/kg have been given. Dosage in renal impairment Reduced dosage is only required in severe renal 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.





Indications Rheumatoid arthritis, osteo-arthritis, low back pain, musculo-skeletal disorders such as fibrositis and other soft-

disorders such as fibrositis and other soft-tissue inflammatory conditions.

Dosage Adults only: 600mg daily in divided doses. Current research suggests that it is not necessary to modify the dosage of Surgam in the elderly or in cases of mild to moderate renal impairment. In severe renal impairment, it is suggested that the dosage should be reduced to 200mg twice daily.

Contra-Indications Active peptic ulceration, hypersensitivity.

Precautions Use with care in patients with a history of peptic ulceration, severe renal or hepatic insufficiency, asthma, sensitivity to aspirin or other non-steroidal anti-inflammatory agents (NSAIs), incipient or actual congestive heart failure. Modification of dosage may be necessary in concurent therapy with highly protein-bound drugs, e.g. anticoagulants, sulphonamides and hypoglycaemic agents. Safety in pregnancy and

lactation has not been established and, in common with other NSAIs, administration during the first trimester should be avoided. Side effects Gastro-intestinal upset, headache, drowsiness and skin rash have occasionally been reported.

Overdosage Supportive and symptomatic therapy.

Presentation and packaging
Surgam 300mg tablets in bottles of 60.
Surgam 200mg tablets in bottles of 100.
Legal Category POM.

Basic NHS prices
Surgam 300mg £14.85 per pack of 60.
Surgam 200mg £15.76 per pack of 100.
Product licence holder and numbers
Roussef Laboratories Ltd., Broadwater Park, North Orbital Road, Uxbridge, Middlesex
UB9 5HP.Surgam 300mg 0109/0109
Surgam 200mg 0109/0108
Date of preparation February 1984.

ROUSSEL 35 Boulevard des Invalides, 75007 Paris, Cédex 07, France



ABC of 1 to 7

H B VALMAN

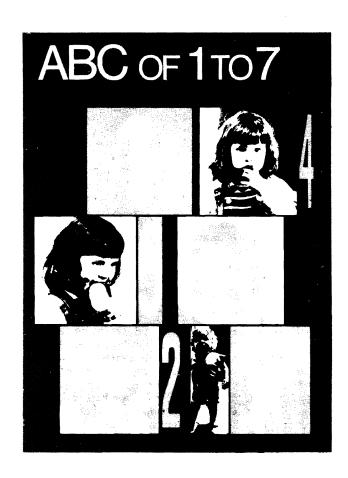
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Postgrad Med J 1983; 59: 273

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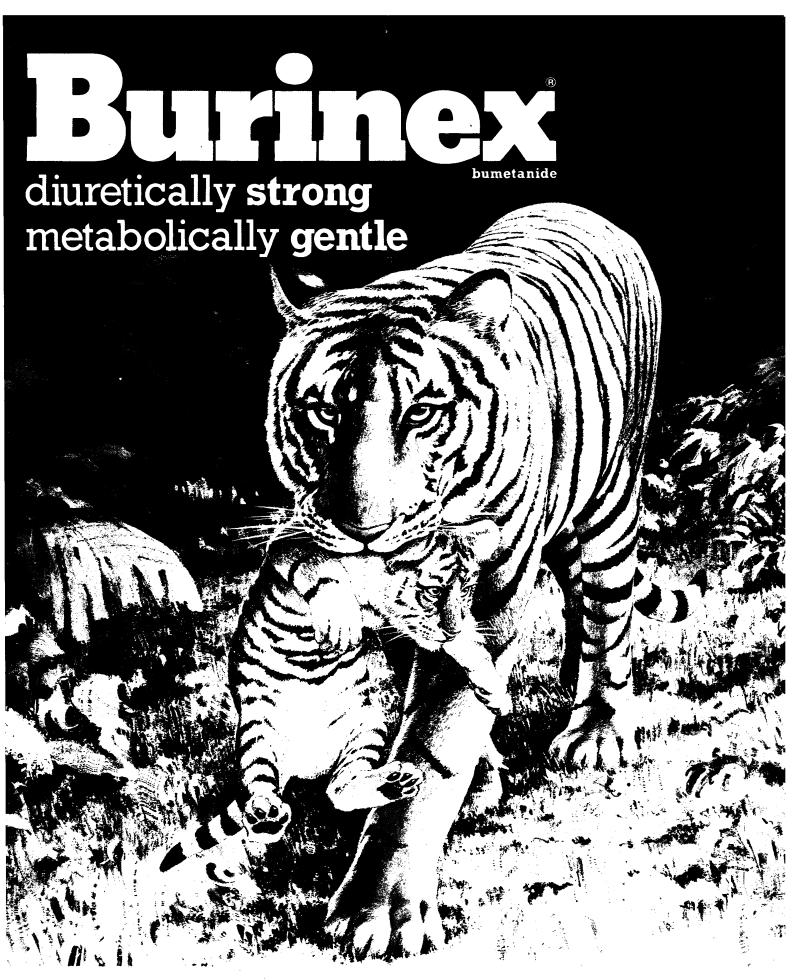
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OCIUM TRADEMARK loperamide

Oral rehydration and the case for imodium In acute diarrhoea it is often essential to treat intravioration his administration finishe intravioration has administration for a definition has been administration for a definition for a definition has been administration for a definition fo in acute diarmoea it is often essential to treat administering fluids intraved administering fluids therand administering therand therapy administering the mouth of his month denydration by administering fluids intravedenydration by administering fluids intravedenydration by mouth. Oral rehydration of characteristic and mouth of characteristic nously of by mouth. Ural renydration therapy mouth. Ural renydration therapy absorption. Will significantly improve the rate of absorption of out it can hardly reduce excessive secretion of out it can hardly reduce. Imodium has been shown to reduce the ex-Impalum has been shown to reduce the electric that the intestinal lument the intestinal lument that the control of the control cessive secretion into the intestinal lumen, the cessive secretion into the intestinal lumen, the cessive secretion into the intestinal the reading the can evading the line and evading the intestinal lumen. July and electrolytes. In fact. Imodium can fluids and expedite the effect of rehydration fluids and expedite the effect of recovery

patient's recovery.

the drug discovery company



Prescribing Information:- Indications Oedema of renal, cardiac or hepatic origin. Dosage Most patients require Img Burinex daily given as morning or evening dose. In refractory cases dosage can be increased to achieve the desired response. For high dose treatment 5mg Burinex should be given initially and increased by 5 mg steps at 12-24 hour intervals until desired response is achieved. Contra-indications, Precautions and Side Effects Contra-indicated in hepatic coma, severe electrolyte depletion and severe progressive

renal failure. Hypovolaemia and circulatory collapse may follow inappropriately excessive diuresis. Electrolyte disturbances resulting in digitalis toxicity may occur. Concurrent antihypertensive or antidiabetic therapy may require adjustment. Caution should be exercised in first trimester of pregnancy. Side effects such as skin rashes, muscular cramps, rises in serum uric acid and thrombocytopenia may rarely occur. Product Licence Number: 1 mg tablets 0043/0021.

Basic N.H.S. Price: £5.43 per 100.



Leo Laboratories Limited Longwick Road, Princes Risborough, Aylesbury, Bucks.

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