## A CONTRACTOR OF THE RELEASE OF THE R

# The first oral anti-fungal effective against all pathogenic fungi

vaginal candidosis:	all other superficial and systemic fungal infections:
2 tablets once daily (with food) for 5 days	tablet daily     (with food)     until complete sympto- matic and mycological     cure is obtained

#### PRESCRIBING INFORMATION

Presentation: white, flat, half scored uncoated tablets marked. "Janssen" on one sight and TuZO0 on the reverse Each fablet contains 200 mg leterocaracity. **Class:** Nicoral is an orally active antimyogate for the treatment in adults of vaginal candidosis, superficial and systemic mycoses including dermatophyte and yeast infections of the skin, har and nails, yeast infections of the mouth and GL tract. Also maintenance treatment of systemic mygoses and severe local infections where previous topical treatment has failed. **Side-effects, precautions, contra-indibations:** contra-indicated in pregnancy. For maximal absorption Naizeral, should be taken with meels. The use of agents which reduce gastine acidity (anti-cholinergic drugs, antacids, I/2 blockers) should be taken with meels. The use of agents which reduce gastine acidity (anti-cholinergic drugs, antacids, I/2 blockers) should be taken with meels. The use of agents which reduce gastine acidity (anti-cholinergic drugs, antacids, I/2 blockers) should be avoided and, if indicated, such drugs should be taken with more tests than two hours after Naizeral. Naizera, etc. in as hybridache and printis 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 indisoyncratic adverse reaction to the drug. This should be bornen 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.

Not all indications are as yet approved in all countries.

Janssen Pharmaceutica B-2340 Beerse, Belgium



#### HISTORY OF THE BRITISH MEDICAL **ASSOCIATION**

Volume II 1932-1981

by

**ELSTON GREY-TURNER** CBE, MC, TD, MD, MRCS, LRCP (Secretary, BMA, 1976-79)

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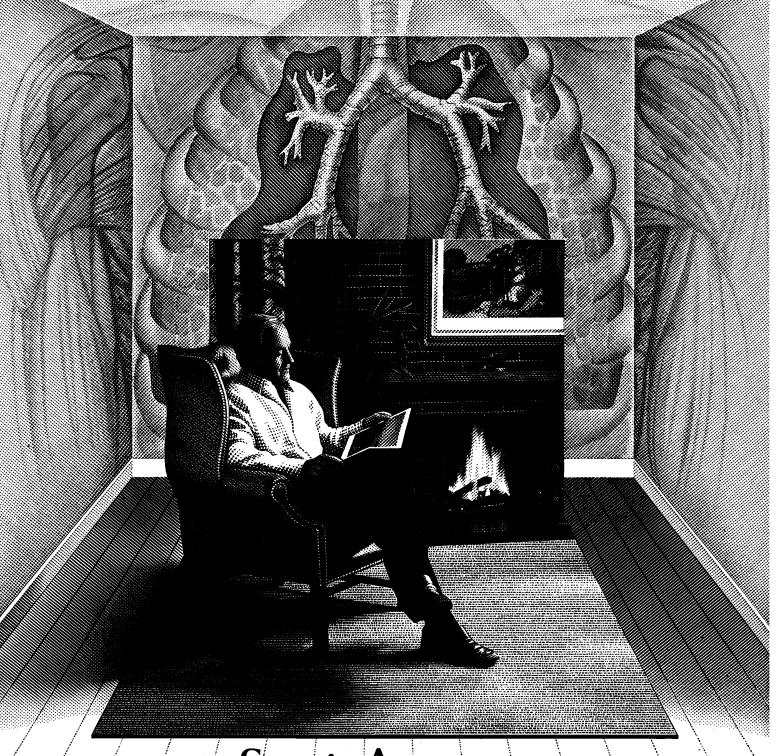
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#### Septrin Assurance

#### Prescribing Information

Indications Sensitive bacterial infections of the lower respiratory, urinary and genital tracts, sinusitis, otitis media, skin infections, septicaemia, typhoid and paratyphoid fevers, and other infections caused by sensitive organisms.

Dosage Septrin Forte Tablets. Adults and children over 12 years: 1 forte tablet twice daily. Maximum dosage for particularly severe infections 1½ forte tablets twice daily. In acute infections Septrin should be given for a minimum of five days or until the patient has been symptom-free for two days. Contra-indications Septrin is contra-indicated in

patients with marked liver parenchymal damage, blood dyscrasias or severe renal insufficiency. Septrin should not be given to patients hypersensitive to sulphonamides, trimethoprim or co-trimoxazole; should not be given during pregnancy or to neonates. Precautions In renal impairment a reduced dosage is indicated and an adequate urinary output should be maintained. Regular blood counts are necessary whenever long-term therapy is used. Caution is advised in patients with folate deficiency. Care should be taken when giving Septrin to patients receiving oral anticoagulants of the coumarin group, pyrimethamine or sulphonylureas.

Adverse Reactions Occasionally, nausea, vomiting, glossitis and skin rashes may occur with normal doses and, very rarely, haematological reactions.

Presentation Septrin Forte Tablets each contain

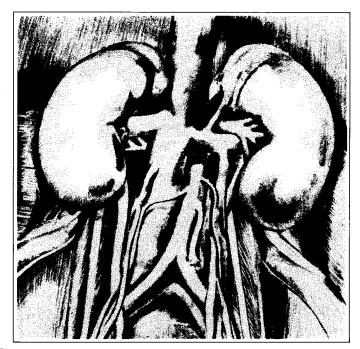
160 mg Trimethoprim BP and 800 mg Sulphamethoxazole BP. PL3/0121.

#### eptrin\* Forte 1b.d.

Further information is available on request. Wellcome Medical Division The Wellcome Foundation Ltd., Crewe, Cheshire



## CALCISORB sodium cellulose phosphate



or the treatment of absorptive hypercalciuria associated with recurrent formation of renal calculi

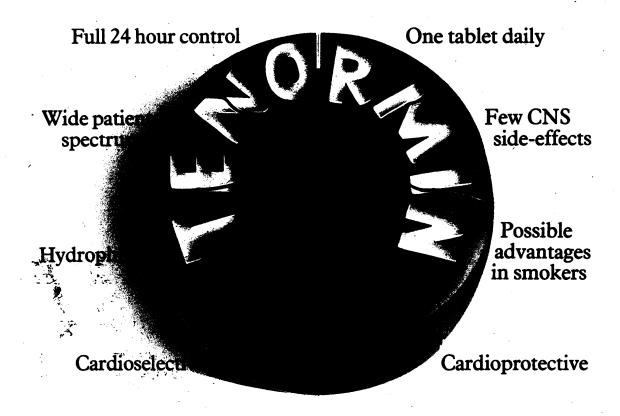
Prescribing information. Composition: Each 5g sachet contains 4.7g sodium cellulose phosphate. Indications: Calcisorb is used to diminish calcium absorption from the diet: 1) in the treatment of hypercalciuria and recurrent formation of renal stones 2) in osteopetrosis 3) as a basis of a test for calcium absorption. Other possible uses are: 1) treatment of hypercalciaemia of infancy 2) treatment of hypercalcaemia csaccidosis 3) treatment of vitamin D intoxication. Dosage and administration: Adults: 15g doily, divided as three 5g doses with meals. Children: 10g daily, divided as three doses with meals. The required dose should be dispersed in water and taken orally. Alternatively the powder may be sprinkled onto food. Side-effects: Side-effects are rare. Isolated cases of diarrhoea have been reported. One patient with mild renal disease, developed a moderate magnesium deficiency. This was readily corrected by halving the dose. Caution: No signs of calcium deficiency have been reported during the continuous use of cellulose phosphate for up to 11 years. This theoretical hazard is particularly relevant to pregnancy, but in view of the absence of data on the effect of cellulose phosphate on calcium levels in pregnant women it is recommended that treatment is discontinued during pregnancy and lactation. Likewise growing children should be prescribed Calcisorb only at the discretion of a senior physician and under his direct supervision. Contra-indications: Renal failure. Congestive heart failure and other conditions in which a low sodium intake is essential. Further Information: Calcisorb should be used in conjunction with a low calcium diet in which dairy products in particular are severely restricted. Basic NHS price: Packs of 100 x 5g sachets: £18.22. Product licence number: 68/5900. Calcisorb; Riker; 3M: trade marks. Further information is available on request.



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Prescribing Notes:

