

Prescribing Information
Uses: 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: Septini Forn. Eiblists, over 12 years, one twice daily. Septini Tibblists. Septini Dispiriblist Eiblists over 12 years, two twice daily, children 6 to 12 years, 10ml Adult twice daily; children 6 to 12 years, 10ml Adult twice daily; children 6 to 12 years, 10ml Adult twice daily; children 6 to 12 years, 10ml Adult twice daily; children 6 to 12 years, 10ml Adult twice daily; children 6 to 12 years, 10ml Adult twice daily; 6 months to 6 years. 5ml Paediatric twice daily; 6 weeks to 6 months, 2.5ml Paediatric twice daily; 6 weeks to 6 months, 2.5ml Paediatric twice daily; 6 months 10 years, 10ml paediatric twice daily; 6 weeks to 6 months, 2.5ml Paediatric twice daily; 6 months 10 years, 10ml paediatric twice daily; 6 weeks to 6 months, 2.5ml Paediatric twice daily; 6 months 10 years, 10ml paediatric twice daily; 6 months 10 years, 10

sml Paediatric twice daily; o weeks to 6 months, 2:ml Paediatric twice daily.

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 or co-trimovable; should not be given during pregnancy or to neonates.

Precautions: In cases of renal impairment a reduced dosage is indicated and an adequate urinary output should be maintained.

in otitis media co-trimoxazole

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 coumanin group, pyrimethamine, sulphonylureas, or

phenytoin.

Warnings and Adverse Effects: Occasionally nausea, vomiting, diarrhoca, glossitis and skin rashes may occur with normal doses and, very rarely, haematological reactions.

Further information is available on request.

Wellcome Medical Division
The Wellcome Foundation Ltd, Crewe, Cheshire.



Wellcome

Presentations:

Product Licence PL3 0(2) Formulation 160mg Francethorram BP 800mg Sulphamethorram ole HP 80mg TAP 100mg SMX 80mg TAP 400mg SMX 80mg TAP 400mg SMX 80mg TAP 400mg SMX 90mg TAP 200mg SMX in 5ml 200mg SMX 10mg SMX 10mg SMX 10mg SMX 10mg SMX Septrin Forte Tablet Septrin Tablets Septrin Dispersible Tablets Septran Adult Suspension Septim Paediatric Suspension Septim Paediatric Tablets PI.3 5222 P1.3 0108

I. Willner, M.W., Dull, T.A., MacDonald, H. 1977. Corr. Chemother, 1, 125. 65 children with acute otitis media involving. Brop. pneumonia. received Septim therapy for an average of 9.3 days. 64. 98.5%, enjoyed complete clearance of clinical symptoms with no evidence of antimicrobial failure.

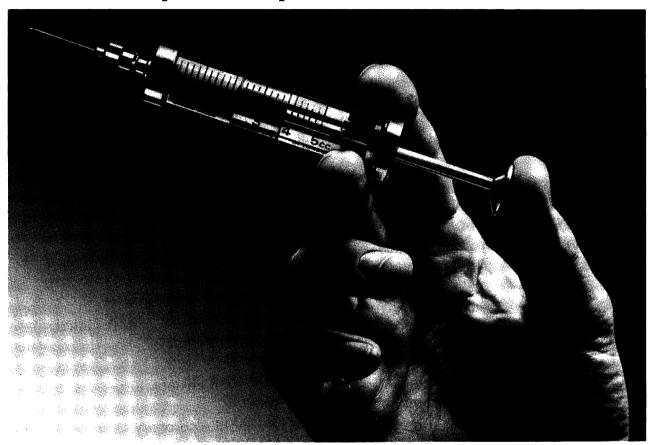
Basic NHS Cust

(1.90 for to

£2.25 tor 20 £2.42 tor 20 £3.22 for 100ml £2.00 for 100ml £0.69 for 20

When a patient comes to you for a jab, maybe you should recommend Flora.

Perhaps the best time to start your patients thinking more seriously about their health is when they come to you about some more routine preventive steps.



It is now widely accepted that changes in diet and lifestyle can dramatically decrease the risk of Coronary Heart Disease developing.

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Flora is high in essential polyunsaturated fats (cis-cis linoleic), because it is made with pure sunflower oil.

Whenever an opportunity arises to recommend ways in which your patients could reduce the CHD risk factors, recommend they change to Flora, too.

Flora. High in essential polyunsaturates.

'Inderal': abridged prescribing information. Presentation: Tablets containing 10, 40, 80, or 160 mg propranolol hydrochloride. **Dosage**: <u>Hypertension</u> 80 mg b.d., increasing weekly. Usual range 160-320 mg daily. Angina 40 mg b.d. or f.i.d., increasing weekly. Usual range 120-240 mg daily. Post myocardial infarction Starting 5-21 days post myocardial infarction, 40 mg q.i.d. for 2-3 days, then 80 mg b.d. Contraindications: Heart block. Bronchospasm. Prolonged fasting. Metabolic acidosis. Precautions: Untreated cardiac failure. Bradycardia. Bronchospasm. Prolonged fasting. Metabolic acidosis. **Precautions**: Untreated cardiac failure. Bradycardia. Modification of tachycardia of hypoglycaemia. Transferance from or discontinuance of clonldine. Prescription of Class I antidysrhythmic agents. Co-administration of verapamil. Anaesthesia. Pregnancy. **Adverse reactions**: Cold extremities, nausea, insomnia, lassitude and diarrhoea are usually transient. Isolated cases of paraesthesia of the hands; rashes and dry eyes have been reported with beta-blockers. Consider discontinuance if they occur. Beta-blockers should be withdrawn gradually. **Overdosage**: See data sheet. **Pack sizes and basic NHS costs** 10 mg 100 :£1.18.1000 :£1.180.40 mg 100 :£3.97.1000 :£39.70.80 mg 60 :£3.78. 500 :£31.48.160 mg 60 :£7.56.250 :£31.48. **PL Nos** 0029/5063, 0029/5064, 0029/5065, 0029/0103. 'Inderal' is a trademark for Propranolol Hydrochlonde BP. Full prescribing information is available from: Imperial Chemical Industries PLC, Pharmaceuticals Division, Alderley House, Alderley Park, Macclesfield, Cheshire SK10 4TF.



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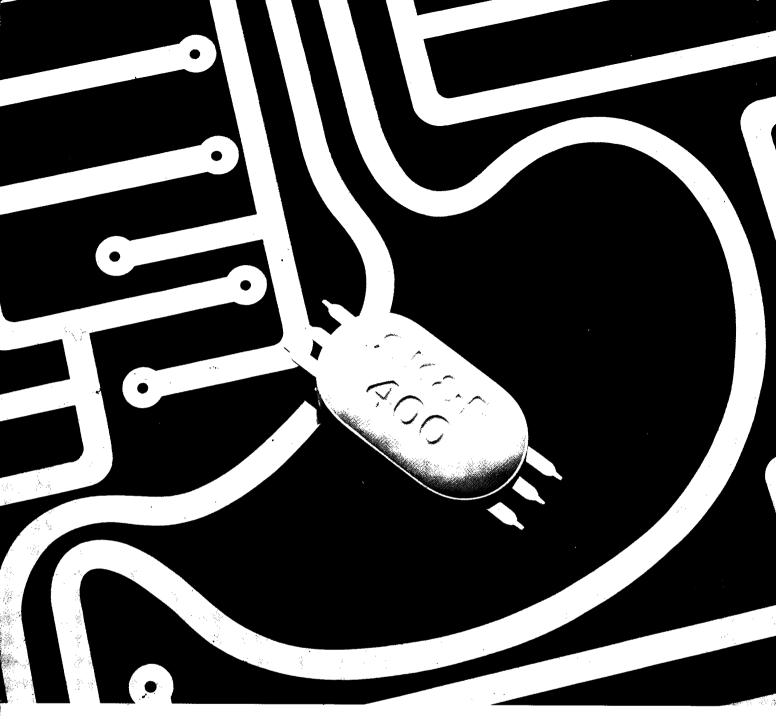
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Preliminary enquiries and requests for further details and application forms should be directed to: Dr H. J. Wright, Division of General Practice, St James's Hospital, Leeds (telephone 0532 433144, extension 5654).



Gastro-technology

Tagamet acid controlled

Prescribing Information. Presentations 'Tagamet' Tablets, PL 0002/0092, each containing 400 mg cimetidine. 56, £16.61. 'Tagamet' Tablets, PL 0002/0063, each containing 200 mg cimetidine. 500, £74.15. 'Tagamet' Syrup, PL 0002/0073, containing 200 mg cimetidine per 5 ml. 500 ml, £20.43. Indications Duodenal ulcer, benign gastric ulcer, recurrent and stomal ulceration, oesophageal reflux disease. Other conditions where reduction of gastric acid is beneficial: prophylaxis of stress-induced gastrointestinal haemorrhage and of acid aspiration (Mendelson's) syndrome; malabsorption and fluid loss in short bowel syndrome. Zollinger-Ellison syndrome. Dosage Usual dosage: Adults. Duodenal ulcer, 400 mg b.d. with breakfast and at bedtime, or 200 mg t.d.s. with meals and 400 mg at bedtime (1.0 g/day) for at least 4 weeks. To prevent relapse, 400 mg at bedtime or 400 mg morning and at bedtime (1.0 g/day) for at least 6 weeks. Desophageal reflux disease, 400 mg at bedtime (1.6 g/day) for at least 6 weeks. Desophageal reflux disease, 400 mg at bedtime (1.6 g/day) for at least 6 weeks. Prophylaxis of stress-induced gastrointestinal haemorrhage, up to 2 g a day, divided, to maintain intragastric pH above 4. Prophylaxis of acid aspiratiom syndrome, 400 mg 90-120 mins before induction of general anaesthesia. 400 mg at start of labour then 200 mg 2-hourly as necessary, suggested maximum 1.6 g. Do not use 'Tagamet' syrup. Zollinger-Ellison syndrome, up to 2 g a day, divided. Recurrent and stomal ulceration and short bowel syndrome, 200 mg t.d.s. and 400 mg at bedtime (1.0 g/day). N.B. For full dosage instructions see Data Sheet. Cautions Impaired renal function: reduce dosage (see Data Sheet). Potentiation of oral anticoagulants, phenytoin and theophylline (see Data Sheet). Prolonged treatment: observe patients periodically. Exclude malignancy in gastric ulcer. Care in patients with compromised bone marrow (see Data Sheet). Avoid during pregnancy and lacatation. Adverse reactions Diarrhoea, dizziness, rash, tiredness. Rarely, mild g

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