#### When gut spasm has 'em

Even if your patients persist with the diet you recommend, it may not be enough to control the pain and spasm of irritable bowel syndrome.

If that is the case, it's a good case for new Merbentyl 20.

A 28-day t.d.s. course of this new presentation of an established antispasmodic should resolve the problem.

So when diet alone just won't do, remember to get it right by writing Merbentyl 20.

#### PRESCRIBING INFORMATION

PRESENTATION: White, biconvex, oval tablets, stamped Merbentyl 20 containing Dicyclomine Hydrochloride BP 20 mg. USES: Merbentyl is a smooth muscle antispasmodic primarily indicated for the treatment of functional conditions involving smooth muscle spasm of the gastro-intestinal tract.

DOSAGE & ADMINISTRATION: Adults and children over 12 years: One tablet (20 mg) three times daily before or after meals.

CONTRA-INDICATIONS, WARNINGS, ETC: Known idiosyncrasy to Dicyclomine Hydrochloride BP.

PRECAUTIONS: **Products** containing dicyclomine hydrochloride should be used with caution in any patient with or suspected of having glaucoma or prostatic hypertrophy. Use with care in patients with hiatus hernia associated with reflux oesophagitis because anticholinergic drugs may aggravate the condition. Since the risk of teratogenicity cannot be excluded with absolute certainty for any product, the drug should be used during pregnancy only if clearly needed.

It is not known whether dicyclomine is secreted in human milk. Because many drugs are excreted in human milk, caution should be exercised when dicyclomine is administered to a nursing woman.

SIDE-EFFECTS: Side-effects seldom occur with Merbentyl. However, in susceptible individuals, dry mouth, thirst and dizziness may occur. On rare occasions, fatigue, sedation, blurred vision, rash, constipation, anorexia, nausea and vomiting, headache and dysuria have also been reported.

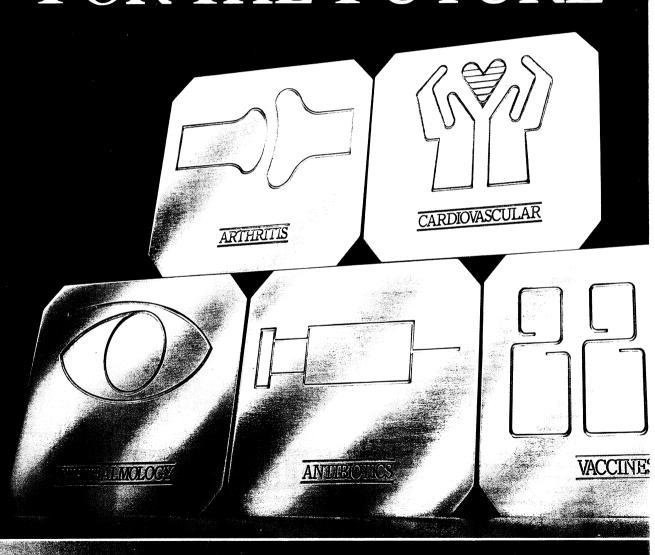
PHARMACEUTICAL PRECAUTIONS: None. LEGAL CATEGORY: POM PACKAGE QUANTITIES: Packs of 84 tablets. FURTHER INFORMATION: Nil. PRODUCT LICENCE NUMBERS: PL 4425/ 0081, PA41/5/1. BASIC NHS PRICE: 84 tablets £4.89 (Oct. 1986). NAME AND ADDRESS OF LICENCE HOLDER: Merrell Dow Pharmaceuticals Limited, Stana Place, Fairfield Avenue, Staines, Middlesex TW18 4SX. TRADEMARKS: Merrell, Dow, Merbentyl.

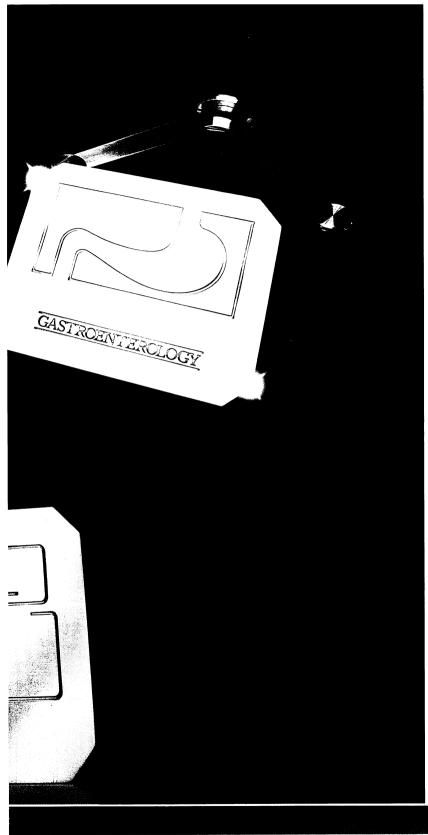
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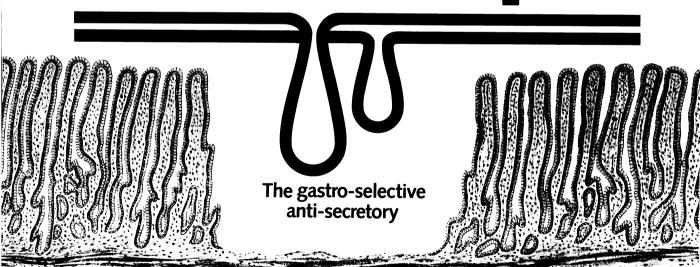
- rely on acid reduction alone
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- profoundly affect intragastric pH

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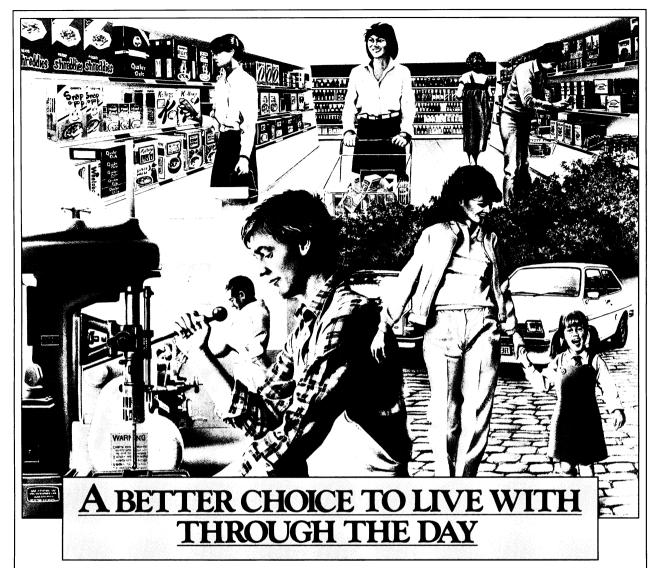
Twice daily
GASTRO I SELECTIVE
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Prescribing Information: Presentation: White tablets each containing 50 mg of prenzepine dhydrochlonde scored on one face with "G" on one side of the score; and "50" on the other The obverse is impressed with the symbol **B** Uses: Gastrozepin is indicated in the treatment of gastric and duodenal ulcers. Dosage: 50 mg at bedtime and in the morning before meals in in severe cases the total daily dose may be increased to 150 mg in divided doses. Continuous therapy may be recommended for up to three months. Contra-indications, Warnings etc: interaction with sympathommentics and monoamme coxidase inhibitors and Gastrozepin is a theoretical possibility. Gastrozepin is not recommended during pregnancy although in animal

experiments no tradoperic effects were noted Breast milk concentration after therapeutic doses is unlikely to affect the infant Side effects occasionally transitory dry mouth and accommodation difficulty may occur freatment of overdosage entirely symptomatic There is no specific antidote Basic NHS price: 50 mg tablets, 600 220 50 Product Licence No.: 50 mg tablets, PLOO34 0260





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References (1) Somerville KW et al. British Medical Journal 1985;291:866. (2) Ruddell WSJ et al. Gut 1980:21:885-889, (3) Farthing MGI et al. British Medical Iournal 1979;2:822-824, (4) Rhodes IM, Journal of Clinical & Hospital Pharmacy 1983;8:219-232. (5) Gaucher P and Champignuelle B. Revue Française de Gastroenterologie 1983;193:35-39. (6) Barr WH et al Medical College of Virginia/Virginia Commonwealth University, FDA bioavailability submission document October 1981. (7) Lee DAH et al. Gut 1980;21:215-218 (8) MIMS October 1985

Prescribing Information: Presentation White odourless acrosol foam containing hydrocortisone acetate PhEur 10%. Uses Anti-inflammatory corticosteroid therapy for the topical treatment of ulcerative colitis, proctosigmoiditis and granular proctitis. Dosage and administration One applicatorful inserted into the rectum once or twice daily for two or three weeks and every second day thereafter. Shake can vigorously before use, (illustrated instructions are enclosed with every pack). Satisfactory response usually occurs within tive to seven days. Contra-indications, warnings etc. Local contra-indications to the use of intrarectal steroids include obstruction, absess, perforation, peritoritis, fresh intestinal anastomoses and extensive fistulae. General precautions common to all corticosteroid therapy should be observed during treatment with Coliform. Treatment should be administered with caution in patients with severe ulcerative disease because of their predisposition to perforation of the bowel wall. Safety during pregnancy has not been fully established. Pharmaceutical precautions Pressurized container. Protect from sunlight and do not expose to temperatures above 50°C. Do not pierce or burn even after use. Do not refrigerate. Shake vivorously before use. Keep our of reach of children. For external use only Legal category POM. Package quantities Aerosol canister containing 25g. (approx. 14 applications) plus a plastic applicator and illustrated leafler. Basic NHS cost 25g plus applicator, £7.25. Further Information One applicatorful of Coliform provides a dose of approximately 125mg of hydrocortisone acetate, similar to that used in a retention enema, for the treatment of ulcerative colius, sigmoiditis and procitits. Product Licence No. 0036/0021. Further information is available on request Statford-Miller Ltd., Professional Relations Division, Harfield, Herts. AL 10 0N2

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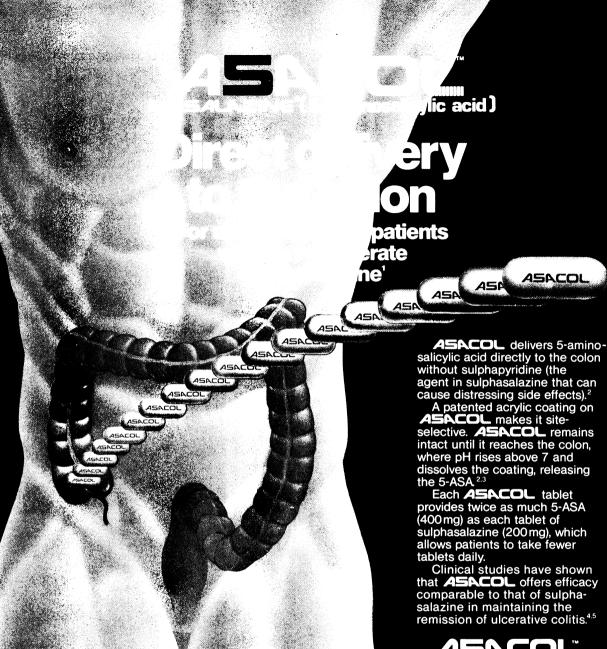
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Dew M.J. Harries A.D. Evans B.K. et al. Treatment of ulcerative colitis with oral 5-aminosalicylic acid in patients unable to take sulphasalazine. *Lancet*, 1983; <u>u</u> 801.

- unable to take sulphasalazine. Lancef. 1983. II. 801. 2 Dew M.J. Hughes P.J. Lee M.G. et al. An oral preparation to release drugs in the human colon. Br. J. Clin. Pharmacol. 1982; 18, 405-408.

  3. Dew M.J. Hyder R.E.J. Evans N. et al. Colonic release of 5-aminosalicylic acid from an oral preparation in active ulcerative collis. Br. J. Clin. Pharmacol. 1983; 15: 185-187.

  4. Dew M.J. Hughes P.J. Harries A.D. et al. Maintenance of remission in ulcerative collis with oral preparation of 5-aminosalicylic acid. Br. Med. J., 1982; 285-1012-1014.

  5. Dew M.J. Harries A.D. Evans N. et al. Maintenance of remission in ulcerative collis with 5-aminosalicylic acid in high doses by mouth. Br. Med. J., 1983; 287-23-24.

#### Mesalazine is the British Approved Name for 5-aminosalicylic acid.

#### ABBREVIATED PRESCRIBING INFORMATION

PRESENTATION
Red tablets containing 400mg of mesalazine (5-aminosalicylic acid) coated for release in the terminal ileum and colon

For the maintenance of remission of ulcerative colitis in patients who cannot tolerate sulphasalazine

DOSAGE AND ADMINISTRATION

Adults: 3 to 6 tablets daily in divided doses
There is no dose recommendation for children

#### CONTRA-INDICATIONS, WARNINGS, ETC.

Contra-indications
Contra-indications a history of sensitivity to salicylates
Children under 2 years of age

Precautions

Renal disorder Mesalazine is excreted rapidly by the kidney marrily as its metabolite. N-acetyl 5-aminosalicytic acid. In rats large doses of mesalazine injected intravenously produce tubular and glomerular loscidy. Although no renal loscidy has been reported in patients taking Asacol. it is not recommended in patients with renal impairment and caution should be exercised in patients with a raised blood urea or proteinuria.

salicylic acid directly to the colon without sulphapyridine (the agent in sulphasalazine that can

### **Direct Delivery to the Colon**

#### Asacol should not be given with lactulose or similar prepara-tions which lower stool pH and may prevent release of

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PL: 0424/0032

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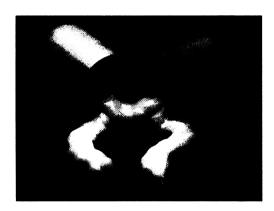
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- Recent ultrasound studies show a consistent inhibitory effect of topical peppermint oil on colon motility and symptomatic improvement of irritable bowel patients given peppermint oil.3

#### References:

- 1. Rees WDW, Evans BK, Rhodes J: Treating irritable bowel syndrome with peppermint oil. *Br Med J* 2:835-836, 1979.
- 2. Somerville KW, Richmond CR, Bell GD: Delayed release peppermint oil capsules (Colpermin) for the spastic colon syndrome: A pharmacokinetic study. Proceedings of the British Pharmacological Society, Cambridge, April 1983. Br J Clin Pharmacol, to be published.
- 3. Taylor BA, Duthie HL, Oliveira RB, et al: Ultrasound used to measure the response of colonic motility to essential oils. Proceedings of *The International Motility Symposium* Aix-en-Provence, France, September 1983, to be published.

(enteric-coated peppermint oil) CAPSULES

#### PRESCRIBING INFORMATION

Presentation: Enteric-coated gelatin capsule. Each contains 0.2 ml standardised peppermint oil B.P. Ph. Eur. Uses: For the treatment of symptoms of discomfort and of abdominal colic and distension experienced by patients with irritable bowel syndrome Dosage and Administration: One capsule three times a day, preferably before meals and taken with a small quantity of water. The capsules should *not* be taken immediately after food. The dose may be increased to two capsules, three times a day when discomfort is more severe. The capsules should be taken until symptoms resolve, usually within one or two weeks. At times when symptoms are more persistent, the capsules can be continued for longer periods of between 2 to 3 months. There is no experience in the use of these capsules in children under the age of 15 years



Contraindications, Warnings, etc. Precautions: The capsule should not be broken or chewed. Patients who already suffer from heartburn, sometimes experience an exacerbation of these symptoms when taking the capsule. Treatment should be discontinued in these patients. Adverse effects: Heartburn, sensitivity reactions to menthol which are rare, and include erythematous skin rash, headache, bradycardia, muscle tremor and ataxia. Product Licence: Pl. 0424/0009. Basic NHS Cost: £10.58 per 100. UK and Foreign Patents pending. Colpermin is a trade mark of Tillotts Laboratories. Further information is available from Tillotts Laboratories Henlow Trading Estate, Henlow, Beds. European Patent No. 0015334.

UK Patent No. 2006011.

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Editorial:

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Original articles:

J.-P. BENHAMOU

mortem cases .....

J.-P. BENHAMOU and F. POTET

CONTENTS

Remarks about esophageal varices and their rupture....

Endoscopic obliteration of esophagogastric varices with

Endoscopic obliteration of esophagogastric varices with

Bucrylate F. II — Histopathological study in 12 post-

B. FABIANI, C. DEGOTT, M.-J. RAMOND, D. VALLA,

Bucrylate®. I — Clinical study in 49 patients......

M.-J. RAMOND, D. VALLA, J.-P. GOTLIB, B. RUEFF and

Nº 8-9

August-September 1986

584

613

Prospective study of patients whose esophageal varices had been obliterated by endoscopic sclerotherapy.....

J. P. VINEL, P. SCHULLER, C. HERVIEU, P. CALÈS, J. CASSI-

J. L. RAOUL, J. F. BRETAGNE, D. MORISOT and J. GAS-

Bacterial meningitis after endoscopic sclerotherapy of esophageal varices: 2 cases......

Cerebral abscess following endoscopic sclerotherapy of

Treatment of bleeding esophageal varices by clip: long-term results in 250 cases.....

Cardiopulmonary arrest after intravenous vitamine K1..

esophageal varices.....

H. LAMOULIATTE, J. C. ARNAL, D. ROUX, A. QUINTON

J. B. PRIOTON, B. FENEYROU, H. MICHEL, P. BORIES.

GARD and P. GAUCHER

L. BERTRAND and F. BLANC

G. PELLETIER, P. ATTALI and O. INK

and J. P. CASTEL

B. CHAMPIGNEULLE, J. M. DOLLET, G. SENG, M. A. BI-

#### GNEUL and J. P. PASCAL DIGESTIVE TRACT AND PANCREAS General review: Editorial: Interferon and liver diseases..... A. Poitrine, S. Chousterman and M. Chousterman Fine needle aspiration cytodiagnosis for cancer of the pancreas (in english) ...... 543 Cerebral infarction following endoscopic obliteration of esophageal varices using Isobutyl-2-Cyanoacrylate: Original articles: report of two cases..... Fine-needle aspiration cytodiagnosis: a simple and safe A. Sée, C. Florent, P. Lamy, V. G. Lévy and M. Bouvry procedure for cancer of the pancreas (in english)..... 545 G. CELLE, V. SAVARINO, E. BIGGI, C. MANSI, P. CEPPA, Letters to the editor: G. R. CICIO and V. ARCURI Carcino-embryogenic antigen in ascites..... Upper digestive tract involvement is frequent in Crohn's E. A. PARIENTE, R. REDIN, B. LUTHIER, M. JALTEL, 549 F. Maitre, A.-M. Kadi and D. H. Trinh H. JOUIN, R. BAUMANN, A. ABBAS, B. DUCLOS, M. WEILL-Infective endocarditis following polypectomy per colono-BOUSSON and J. P. WEILL 609 Contribution of anorectal manometry and ECG to the F. GRÉCO, D. KRAI, A. ZANNETTI, B. BOULAY, Y. ÉTIENNE diagnosis of diabetic autonomic neuropathy...... and J. L. BOURDON D. LEJEUNE, M. MELANGE, Ch. DAUMERIE, M. BUYS-Autoimmune hemolytic anemia associated with ulcerative SCHAERT and R. VANHEUVERZWIJN 609 Myocardial infarction and polyps of the colon...... T. COSTE, R. BENAMOUZIG, P. CASASSUS, V. DUCHATELLE. J. C. Audigier, C. Barthelemy, J. P. Étaix, B. Duran-D. HUC and J. RAUTUREAU TON, H. VERNEYRE and H. FRAISSE Cowden's disease (sixth french case). Efficacy of etretinate on cutaneous and digestive manifestations..... Current trend: J. BRUHIÈRE, J. THIVOLLET and D. JACQUEMIER Bismuth-induced encephalopathies: reappraisal of risks... Answer to the article of Mr. Huguier et al..... 562 611 D. LOISEAU P. LECHAT and R. KISCH Can sclerotherapy of esophageal varices induce cholestasis?..... LIVER AND BILIARY TRACT 611

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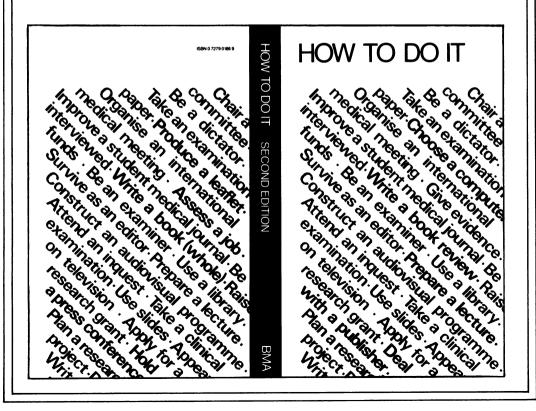
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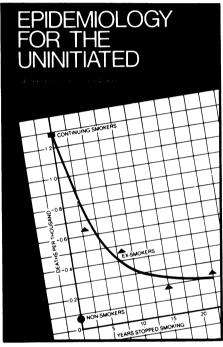
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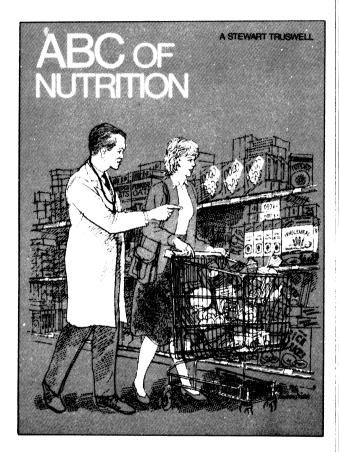
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