

Ease the spasm. Ease the mind.



clidinium bromide and chlordiazepoxide

Clidinium bromide to calm the gut. Chlordiazepoxide to calm the mind.

Indications For the control of hypersecretion, hypermotility and emotional factors associated with gastro-intestinal disorders, such as nervous dyspepsia, peptic ulcer, cardiospasm, pylorospasm, nervous or

Dosage 1 or 2 tablets three or four times daily. In elderly patients, it is recommended that the initial dose be I tablet twice daily.

Contra-indications Because of its anticholine effects, Libraxin should not be given to patients from glaucoma or prostatic enlargement.

Precautions Patients should avoid alcohol while under treatment with Libraxin, since the individual

ROCHE

response cannot be foreseen. Patients' reactions (driving ability, operation of machinery, etc.) may be modified to a varying extent, depending on dosage and individual susceptibility. The established medical principle of prescribing medicaments in early pregnancy only when absolutely indicated should be observed.

Side-effects Side-effects are infrequent and are controlled by reduction of dosage. The

drowsiness, muscle weakness, dryness of the mouth, blurring of vision, constipation and hesitancy of micturition.

Presentation Libraxin tablets containing 5mg chlordiazepoxide and 2.5mg clidinium bromide in packings of 100 and 500.

Basic NHS Cost 1 tablet 3 times daily 7.4p/day ex 500 pack.

Licence Number 0031/5024

Licence Holder Roche Products Limited, PO Box 8 Welwyn Garden City, Hertfordshire AL7 3AY Libraxin is a trade mark

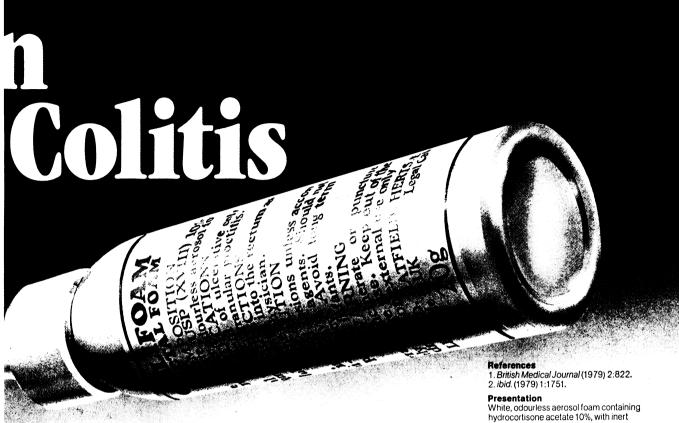
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Colifoam is a unique therapy for ulcerative colitis, being a topical anti-inflammatory with exceptional benefits over the rectal enema in terms of simplicity and convenience.

Gamma photography studies have shown that a single dose of Colifoam remains in contact with the rectal mucosa for several hours. In one of these studies the foam was seen to reach the sigmoid colon in most patients. The second study, using a different protocol which included healthy subjects, did not confirm this finding but the authors concluded:

"Unquestionably, however, the foam is more comfortable and easier to retain



than a retention enema, and since the patient need not be immobilised, the foam obviously has a place in outpatient practice for patients with proctitis and distal ulcerative colitis."

Colifoam: hydrocortisone acetate foam supplied in a metered dose dispenser, delivering approximately 5 ml. of Colifoam rectal foam containing 10% hydrocortisone

Colifoam hydrocortisone acetate foam

comfort and convenience in ulcerative colitis

propellants.

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 in each pack). One applicatorful of Colifoam provides a dose of approximately 90-110mg of hydrocortisone, similar to that used in a retention enema for the treatment of ulcerative colitis, sigmoiditis and proctitis. Satisfactory response usually occurs within five to seven days.

Contra-indications and Warnings etc.

Local contra-indications to the use of intrarectal steroids include obstruction, abscess, perforation, peritonitis, fresh intestinal anastomoses and extensive fistulas. General precautions common to all corticosteroid therapy should be observed during treatment with Colifoam. Treatment should be administered with caution in patients with severe ulcerative diseases because of their predisposition to perforation of the bowel wall. Safety during pregnancy has not been fully established.

Pharmaceutical Precautions

Do not refrigerate, incinerate or puncture the aerosol can. Shake vigorously before use. Keep out of reach of children.

Package Quantities

Aerosol canister containing 20g (14 applications) plus a plastic applicator and illustrated leaflet.

Basic NHS Cost

Product Licence No. 0036/0021

Further information is available on request

Stafford-Miller Limited,

Professional Relations Division, Hatfield. Herts. AL10 0NZ.





Maxolon protects th over-long exposure to g normal peristalsis and ga action contrasts with that

By restoring the stom symptoms described by pain, heartburn and d

treated and their recurrence To the patient. Mass convenient therapy to reantacid prescriptions.

scribing Information

Indications
Dyspepsia, heartburn and flatulence
associated with the following conditions
e.g., Reflux oesophagitis, Gastritis, Hiatus
hernia, Peptic ulcer.

Adult Dosage (oral)
Adults 10mg
1 tablet or 10ml syrup 3 times a day.
Young adults (15-20 years) 5-10mg
½1 tablet or 5-10ml syrup 3 times a day.
commencing at the lower dosage.

Note: Total daily dosage of Maxolon, especially for children and young adults should not normally exceed 0.5mg/kg

Side-effects and Precautions
There are no absolute contra-indications
to the use of Maxolon.

Various extra-pyramidal reactions to Maxolon, usually of the dystonic type, have been reported. The incidence of these reactions in children and young adults may be increased if daily dosages higher than 0.5mg/kg body-weight are administered. The majority of reactions occur within 36 hours of starting treatment and the I he majority of reactions occur within 36 hours of starting treatment and the effects usually disappear within 24 hours of withdrawal of the drug. Should treatment of a reaction be required, an anticholinergic anti-Parkinsonian drug e.g. benaptyzine, or a benzodiazepine may be used. Since extrapyramidal symptoms may occur with both

Maxolon and phenothiazines, care should be exercised in the event of both drugs being prescribed concurrently.

Raised serum prolactin levels have been observed during metoclopramide therapy; this effect is similar to that noted with many other compounds.

Maxolon's action on the gastro-intestinal tract is antagonised by anticholinergics. Although animal tests in several mammalian species have shown no teratogenic effects, treatment with Maxolon is not advised during the first trimester of pregnancy

Following operations such as pyloroplasty or gut anastomosis Maxolon therapy should be withheld for three or four days as

vigorous muscular contractions may not help healing.

Availability and NHS Prices Tablets 10mg (£5.84 per 100). Syrup 5mg/5ml (£2.42 for 200ml).

A paediatric liquid presentation and ampoules for injection are also available. Average daily cost of Maxolon tablets (ex. 500 pack) 17p. Prices correct at January 1979. Further information is available on request to the company.

Maxolon (metoclopramide) is a product of

Beecham Research Laboratories,
Brentford, England.
A branch of Beecham Group Limited.

PL 0038/0095 0098 5040 5041

1. Gut. (1969), 10, 678-680. 2. Postgrad. med. J., (1973), 49, (Suppl), 29. 3. Gut. (1974), 15, 462-467. 4. Brit. med. J., (1971), 2, 25

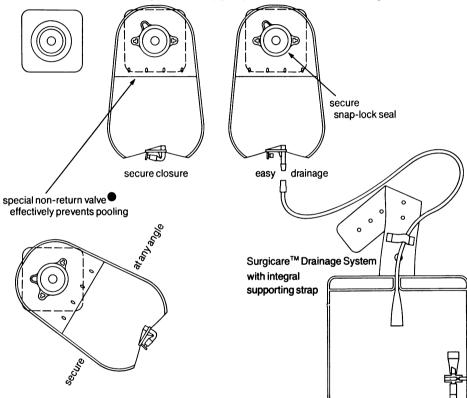
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gastric ulcer and reflux oesophagitis, where acid plays a part.

For many patients it has brought a new standard of pain relief and healing. has been prescribed for an estimated one million patients.

PRESCRIBING INFORMATION

PRESCRIBING INCOMPANIONAL PRESCRIPTION OF THE PRESCRIPTION OF THE

Indications Duodenal ülcer, benign gastric ülcer, reflux oesophagus

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and 440 mg at bedfine. 10g day for the full media
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and 400 mg at bedfine. 10g day, for 4 for 8 weeks.

Cautions

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Sheet. Potentiation of oral anticoagulants (see Data
Sheet. Prolonged treatment observe patients
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symptomatically Avoid during pregnancy and
life from the proposition of the pr

Adverse reactions

Adverse reactions
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states usually in the elderly or very ill, intenstitial
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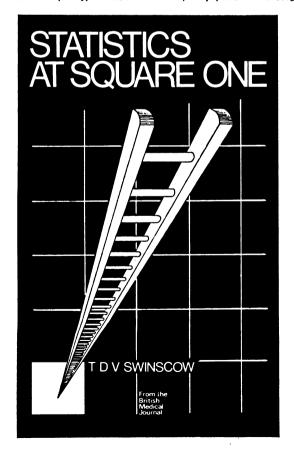
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O'Donnell, Barry, British Medical Journal, 1977, 1, 451.

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Gut August 1980



Pyrogastrone (PL 0071/0138). For the treatment of oesophageal inflammation, erosions and ulcers due to hiatus hernia or other conditions causing gastric reflux and for the relief of heartburn, flatulence and other symptoms associated with reflux oesophagitis. Each tablet contains: carbenoxolone sodium B.P. 20mg, magnesium trisilicate B.P. 60mg, dried aluminium hydroxide gel B.P. 240mg, in a base containing sodium bicarbonate B.P. 210 mg and alginic acid B.P.C. 600 mg. Cartons of 100. Adult Dosage. One to be chewed immediately after meals, three times a day and two to be chewed at bedtime. Basic N.H.S. Cost: One days treatment 56p (5 tablets). Contraindications: Severe cardiac, renal or hepatic failure. Patients on digitalis therapy unless serum electrolyte levels are monitored weekly to detect promptly the development of hypokalaemia. Precautions: Special care should be exercised with patients predisposed to sodium and water retention, potassium loss and hypertension (e.g. the elderly and those with cardiac, renal or hepatic disease) since the carbenoxolone content of Pyrogastrone can induce similar changes. Regular monitoring of weight and blood pressure which should indicate the development of such effects is advisable for all patients. A thiazide diuretic should be administered if oedema or hypertension occurs. (Spironolactone should not be used because it hinders the therapeutic action of carbenoxolone). Potassium loss should be corrected by the administration of oral supplements. No teratogenic effects have been reported with carbenoxolone sodium, but careful consideration should be given before prescribing Pyrogastrone for women who may become pregnant. Pyrogastrone is a registered trade mark. Made under licence from Biorex Laboratories Brit. Pat. No. 1390683. Further information available from:—

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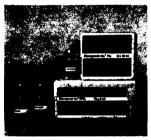
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