## 25000 lipase units



This high lipase formulation of Creon should allow Cystic Fibrosis patients to substantially reduce their daily capsule intake and achieve better compliance.



#### HIGHER DOSE, FEWER CAPSULES, HAPPIER PATIENTS

#### Prescribing Information

**Presentation:** Opaque orange yellow hard gelatin capsules containing brownish coloured enteric coated pellets of pancreatin, equivalent to:

25,000 BP units of lipase

18,000 BP units of amylase

46" BP units of protease

Available in packs of 50. Basic NHS price £19.50

Indication: Pancreatic exocrine insufficiency

**Dosage and Administration:** Adults (including elderly) and children: Initially one capsule with meals, then adjust according to response.

The capsules can be swallowed whole, or for ease of administration they may be opened and the granules taken with fluid or soft food, but without chewing. If the granules are mixed with food it is important that they are taken immediately, otherwise dissolution of the enteric coating may result.

Contra-indications, Warnings, etc. Contra-indications: Substitution with pancreatic enzymes is contra-indicated in the early stages of acute pancreatitis. Warnings: Use in pregnancy: There is inadequate evidence of safety in use during pregnancy. The product is of porcine origin. Rarely cases of hyper-uricosuria and hyper-uricaemia have been reported with very high doses of pancreatin.

Overdosage although not experienced until now, could precipitate meconium ileus equivalent.

Perianal irritation, and rarely, inflammation, could occur when large doses are used.

Product Licence Number: 5~2~ 0006

#### Name and Address of Licence Holder

Kali Chemie Pharma GmbH, Hans-Bockler-Allee 20, 3000, Hannover 1, Germany.

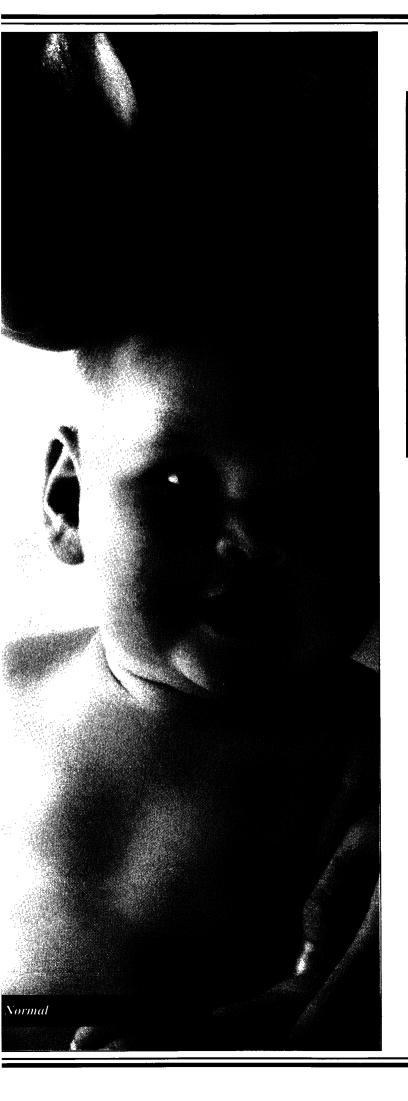
Further information is available from: Duphar Laboratories Limited, Gaters Hill, West End,

Southampton SO3 3[D. Tel: (0703) 472281

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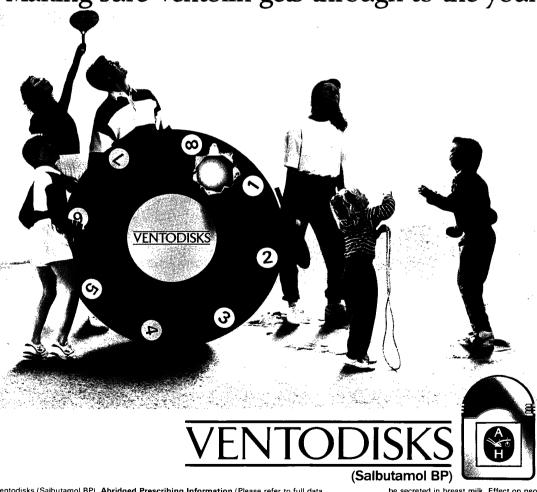
The Babyfeeding Specialists

Breastmilk is the best food for babies. The purpose of infant milk formula is to replace or supplement breastmilk when a mother cannot, or chooses not to, breastfeed. The cost of infant milk formula should be considered before deciding how to feed a baby.

This advertisement has been prepared for the information of the Health Care Professional only.



Making sure Ventolin gets through to the younger generation



Ventodisks (Salbutamol BP). Abridged Prescribing Information (Please refer to full data sheet before prescribing.) Uses: Treatment and prophylaxis of acute and chronic bronchospasm. Dosage and administration: Adults: 400 micrograms as single dose or three to four times daily. Children: 200 micrograms as single dose or three to four times daily. Contra-indications: Threatened abortion during first or second trimester. Hypersensitivity. Precautions: If previously effective dose lasts less than three hours, seek medical advice. Caution in patients with thyrotoxicosis. Avoid use with non-selective beta-blockers. Hypokalaemia may occur, particularly in acute severe asthma. It may be potentiated by xanthine derivatives, steroids, diuretics and hypoxia. Serum potassium levels should be monitored in such situations. Pregnancy: Avoid unnecessary use during early pregnancy. Only consider if expected benefit outweighs possible risks. Lactation: Salbutamol likely to

be secreted in breast milk. Effect on neonate unknown. Balance risks against benefits. Side effects: Mild tremor, headache occur rarely. Very rarely — transient muscle cramps and hypersensitivity reactions. Potentially serious hypokalaemia may result from 82-agonist therapy. Paradoxical bronchospasm could occur – substitute alternative therapy. Presentation and Basic NHS cost: Pack of 14 Ventodisks each containing 8 x 200 micrograms Salbutamol BP (as sulphate) — light blue or 8 x 400 micrograms Salbutamol BP (as sulphate) — dark blue, together with a Ventodin Diskhaler. For inhalation. £7-11 and £12-02. Refill pack of 14 x 8 Ventodisks only. 200 micrograms, £6-54; 400 micrograms, £11-45.



Product licence numbers: Ventodisks 200 micrograms 0045/0134, Ventodisks 400 micrograms 0045/0135.

ALLEN & HANBURYS

PRESCRIBING INFORMATION Presentation TILADE Mint is a metered dose pressurised aerosol inhaler which delivers 56 puffs, each containing 2mg Nedocromil sodium. <u>Indications</u> TILADE Mint is indicated for the preventive treatment of reversible obstructive airways disease, including ashma and ashmatic branchitis. <u>Dosage and Administration</u> Adults and children over 12 years of age: 2 puffs [4mg Nedocromil sodium] to be inhaled twice daily. If necessary, dosage may be increased to 2 puffs four times daily.

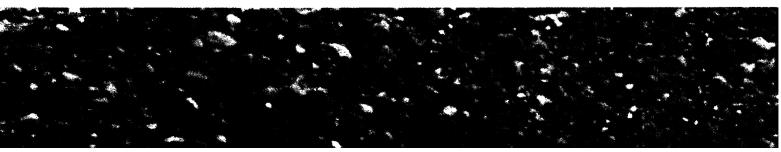
Children under 12 years of age: it is recommended that TILADE Mint should not be used since it is still under investigation in this age group. TILADE Mint is intended for regular daily usage and should not be used for relief of symptoms in an acute attack. Side effects, precautions and contra-indications few side effects have been reported, principally headache and nausea. The mint flavour has been added to mask any bitter taste of the drug. Caution should be exercised in the use of TILADE Mint in pregnant or lactating

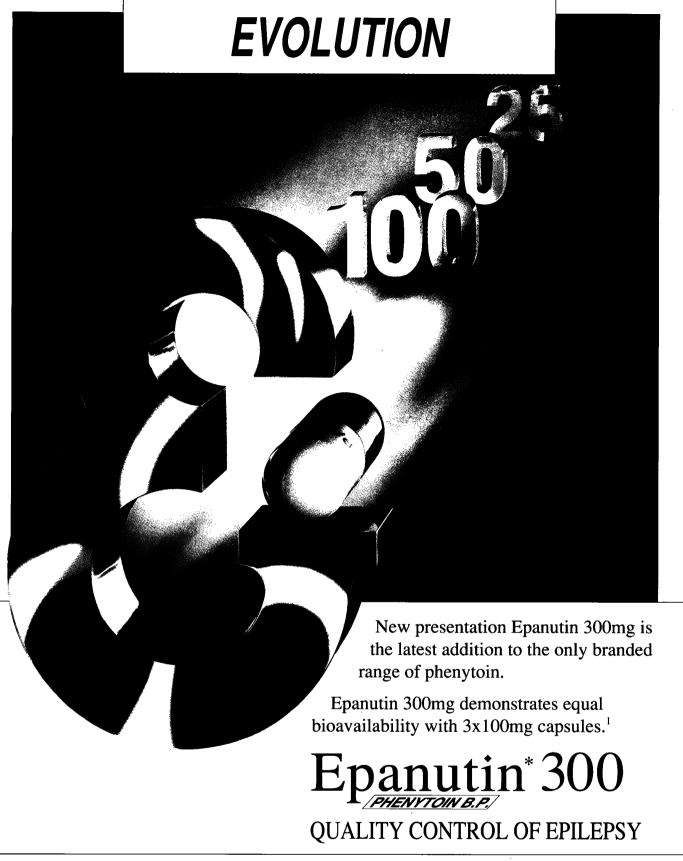
women. There are no specific contraindications. Basic NHS Cost 2 x 56
actuations £18.65. PL Number: 0113/0140.
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Prescribing Information Presentation: Epanutin Capsules, 25mg, 50mg, 100mg or 300mg phenytoin sodium Ph Eur. Epanutin Suspension 30mg/5ml phenytoin BP. Epanutin Infatabs, 50mg phenytoin BP. Indications: Grand mal epilepsy, temporal lobe seizures and certain other convulsive states. Dosage: Usual maintenance dosages: Infants and children: 4-8mg/kg, up to a maximum of 300mg daily. Adults: 200-500mg daily in single or divided doses. Exceptionally, a daily dose outside this range may be indicated. Dosage recommendations are only guidelines and should be titrated for individual patients. Chew Infatabs. Titrate dosage gradually. Plasma level monitoring is advisable. Equal doses of Infatabs and capsules may not give equivalent blood levels. Contra-indications and precautions: Hypersensitivity to hydantoins. Use in pregnancy and lactation. Liver dysfunction. Replacement of or with other anticonvulsant therapy should be gradual. Plasma levels may be altered by other drugs - see literature. Side-effects: Transient GI and CNS disturbances subsiding with continued use. Allergic phenomena (e.g. rash, lupus erythematosus, hepatitis, lymphadenopathy) may occur. Haematological disorders, gingival hypertrophy, hirsutism and excessive motor activity have been reported. Nystagmus

with diplopia and ataxia indicates that dosage should be reduced. **Legal category:** POM. **Product Licence Numbers:** Capsules, 25, 50, 100mg/18/0112, 18/5079, 18/5080, 300mg, 18/0158. Suspension 18/5106. Infatabs 18/0069. **Basic NHS cost:** Capsules, 25mg 500 £9.82; 50mg 500 £10.02; 100mg 500 £12.77;1000 £24.15; 300mg 100 £7.66; Infatabs 100 £5.49. Suspension 500ml £3.56 Further information is available from Parke-Davis Research Laboratories, Lambert Court, Chestnut Avenue, Eastleigh, Hampshire SO5 3ZQ. Telephone: (0703) 620500.

#### References:

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**New formulation replaces Vamin® Infant** 

New Vaminolact® is formulated to closely resemble the natural source of protein for premature and older infants — human milk protein. So it's only natural that it should contain the right balance of essential and non-essential amino acids, like low phenylalanine and added taurine. New sulphite-free Vaminolact, specifically for use in paediatric parenteral regimens — when all that matters are the little things.

FIRST LINE AMINO ACIDS FOR NEONATES AND INFANTS

Kabi Pharmacia

#### ABBREVIATED PRESCRIBING INFORMATION

PRESENTATION: A solution for intravenous nutrition, specifically formulated for paediatric use.

L-Alanine	6. <b>3</b> g
L-Arginine	4.1g
L-Aspartic acid	4.1g
L-Cysteine/cystine	1.0g
L-Glutamic acid	7.1g
Glycine	2.1g
L-Histidine	2.1g
L-Isoleucine	3.1g
L-Leucine	7.0g
L-Lysine	5.6g
L-Methionine	1.3g
L-Phenylalanine	2.7g
L-Proline	5.6g
L-Serine	3.8g
Taurine	0.3g
L-Threonine	3.6g
L·Tryptophan	1.4g
L-Tyrosine	0.5g
L-Valine	3.6g
In each 1000ml	65.39
Osmolality: 510mosmol	per kg
water.	

**Nitrogen per litre:** 9.3g corresponding to 58g of protein.

Energy content per litre: 240kcal

Indications: Vaminolact should be used for the prophylaxis and therapeutic treatment of protein depletion in neonates and infants

depletion in neonates and infants where sufficient enteral nutrition is impossible or impracticable.

DOSAGE AND ADMINISTRATION:
Recommended dosage for infants: the dosage should be increased gradually during the first week of administration to a final daily described to a Sami Assemblact neck. dose of up to 35ml Vaminolact per kg body weight given as a continuous infusion over 24 hours. The duration of infusion should be at least 8 hours.

Recommended dosage for children: The dosage should be increased gradually during the first week of administration to the final dosage ranges indicated below and should be infused over 24 hours

	Dosage* (ml per bodyweight per
Weight (kg)	24 hours)
10	24.0
20	18.5
30	16.0
40	14.5

\*The duration of infusion should be at least 8 hours.

(ml per kg

at least 8 hours.

Caution: The recommended infusion rate should not be exceeded.

Contra-indications, warnings etc: varninolact is contra-indicated in patients with irreversible liver damage and in severe uraemia when dish eich exiliations.

damage and in severe uraemia when dialysis facilities are not available Care must be exercised in the admini-stration of large volume infusion fluids to patients with cardiac insufficiency. Amino acid infusions must also be administered with caution to patients with disturbances in protein metabolism.

In protein metabolism.

Precautions: Hyperkalaemia, hypernatraemia, and acidosis should be corrected prior to commencement of intravenous nutrition. Serum electrolytes, blood glucose levels and acid-base balance should be regularly actionase palantoe should be regularly monitored. Fluid balance should also be monitored since hypertonic dehydration may occur. Amino acid solutions may precipitate acute folate deficiency and folic acid should be

**Side-effects:** Vaminolact is well tolerated. Rarely, nausea may occur. tolerated. Rarely, nausea may occur. As with all hypertonic solutions, thrombophlebitis may occur when peripheral veins are used. Abnormal liver function tests have been observed during intravenous nutrition

Legal category: POM.

Package quantities: Bottles of
100ml and 500ml.

Further information: The manufacturer can be consulted for full information on complete and balanced intravenous nutrition

Product Licence number: 0022/0092.

Product authorisation numbers:

100ml 187/35/1, 500ml 187/35/2

**Product Licence/Authorisation** 

holder: Kabi Pharmacia Limited, Davy Avenue. Kadi rharmacia Limited, Day Avenue, Knowlhill, Milton Keynes, MK5 8PH. Distributed by Cahill May Roberts Limited for Kabi Pharmacia (Ireland), Pharmapark, Chapelizod, Dublin 20. Cost: Vaminolact 100ml £3.80. Vaminolact 500ml £8.70. Pediatrics

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Volume 150 No. 12 1991

The pioneers of pediatric medicine 823

Review

Small-intestinal abnormalities in cystic fibrosis patients
E. Eggermont, K. De Boeck 824

Cardiology

Left ventricular thrombus in a 2-year-old boy with cardiomyopathy: lysis with recombinant tissue-type plasminogen activator O.N.Krogmann, R.v.Kries, S.Rammos, H.H.Kramer, M.Bourgeois 829

Gastroenterology/Hepatology

Documented latent coeliac disease in a child with insulindependent diabetes mellitus C. Catassi, G. Natalini, I. M. Rätsch, O. Gabrielli, G. V. Coppa, P. L. Giorgi 832

Hematology/Oncology

Optic gliomas in children with neurofibromatosis type 1 A. M. Lund, F. Skovby 835

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Cytoplasmic granules in leukaemic cells of the cerebrospinal fluid in a child with non-granular acute lymphocytic leukaemia

T. Hanada, I. Ono, N. Moriyama, K. Koike 839

Transcobalamin II deficiency: case report and review of the literature

Y. Kaikov, L.D. Wadsworth, C.A. Hall, P.C.J. Rogers 841

Immunology/Allergology

Cord serum IgE and early detection of the atopic phenotype: suitable for routine screening? H.S.Varonier, G.C.Lacourt,

A. Assimacopoulos 844

Medical genetics

Cartilage hair hypoplasia in infancy: a misleading chondrodysplasia M. Le Merrer, P. Maroteaux 847

Metabolic diseases

Chronic pancreatitis in a child with glycogen storage disease type 1 M. Kikuchi, K. Hasegawa, I. Handa,

M. Watabe, K. Narisawa, K. Tada 852

Neonatology

Ipratropium bromide for symptomatic preterm infants B. Yuksel, A. Greenough 854

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Mitral valve and tricuspidal valve prolapse in Melnick-Needles syndrome

G. Krüger, K. Schumacher, F. Erfurt, L. Pelz 858

An infant having Potter facies without pulmonary hypoplasia after prolonged oligohydramnios due to a placental mass M.Ohyama 859

Simultaneous epiglottitis siblings B. Zimmer, K. Vogel, D. Schranz 859

Acute caffeine poisoning in a child
P. G. Jorens, J. M. Van Hauwaert

P. G. Jorens, J. M. Van Hauwaert, M. I. Selala, P. J. C. Schepens **860** 

**Abstracts** 

The Summer Meeting of the Scottish Paediatric Society, Aberdeen, 31 May 1991 861

Acknowledgement to our reviewers 865

Contents of Volume 150

Index of key words

Indexed in Current Contents



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More and more patients with uncontrolled epilepsy are able to face the world thanks to the efficacy of SABRIL.

Many have become seizure free for the first time and approximately half can benefit from a > 50% reduction in their seizures.<sup>1</sup>



Controlling seizures, changing lives

#### Abridged Prescribing Information SABRIL Tablets

SABRIL Tablets
Presentation: White, tablets with a breakline marked SABRIL, each containing 500mg vigabatrin.
Uses: Indications: Treatment of epilepsy not controlled by other antiepileptic drugs.
Dosage and Administration: Oral administration once or twice daily added to the patient's current therapeutic regimen.

Adults: Recommended starting dose 2g/day. Increased or decreased in 0.5g or 1.0g increments depending upon clinical response and tolerability. Maximum 4g/day. There is no direct correlation ween plasma concentration and efficacy.

Detween plasma concentration and entracty.

Children: The recommended starting dose in children is 40mg/kg/day increasing to 80-100mg/kg/day depending on response. Convenient recommendations in relation to bodyweight are:

Bodyweight: 10-15kg 0.5 to 1g/day

15-30kg 1.0-1.5g/day

10 -15kg 15-30kg

30-50kg 1.5-3g/day

30-50kg 1.5-3g/day
>50kg 2-4g/day
Infants with West syndrome may require doses of 100mg/kg/day or higher.
Elderly: Consider dose reduction in patients with impaired renal function.
Contra-Indications, Precautions, Warnings etc.

Use in pregnancy and lactation: Contra-Indicated.

Precautions: Abrupt withdrawal may lead to rebound seizures. Caution in patients with history of psychosis or behavioural problems. Caution in elderly patients, particularly creatinine clearance b 60ml/mln. Reduce dose and monitor closely for adverse events.

ings: Vigabatrin causes intramyelinic cedema in the brain white matter tracts of animals but there is no evidence of this in man. However, monitor patients for neurological changes. See the full

Effects on driving ability: Drowsiness has been seen and patients should be warned.

Effects: Are mainly CNS related. Aggression and psychosis have been reported and a previous history of psychosis or a behavioural problem appears to be a predisposing factor. Other reported events: drowsiness and fatigue, dizziness, nervousness, irritability, depression, headache and less commonly confusion, memory disturbance and vision complaints; also weight gain and minor gastrointestinal side-effects. Also in children excitation and agitation. Some patients may experience an increase in seizure frequency with vigabatrin, particularly those with myoclonic seizures have not revealed evidence of neurotoxicity in humans. Lab data do not indicate renal or hepatic toxicity. Decreases in SGOT and SGPT have been observed. Chronic treatment may lead to a slight decrease in haemoglobin.

Drug Interactions: Are unlikely. A gradual reduction of about 20% in plasma phenytoin concentration has been observed. No clinically significant interactions with carbamazepine, phenobarbitone or sodium valproate.

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Package Quentities: Blister strips of 10 in cartons of 100.
Product Licence Number: PL 4425/0098

NHS Price: £46.00

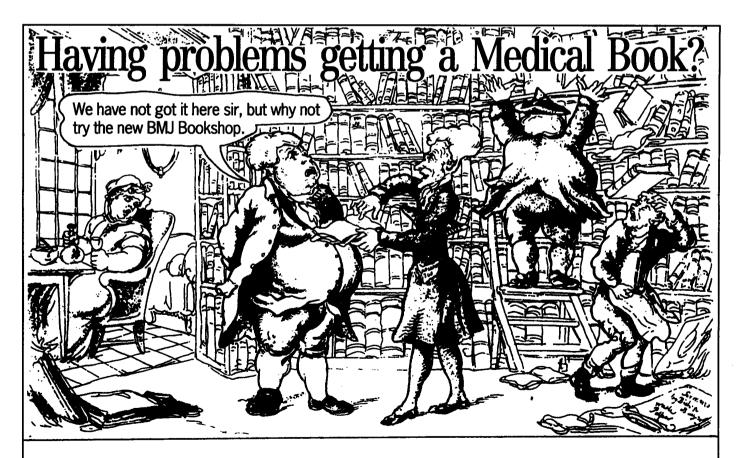
tion: January 1992.

**Date of Preparation:** January You must refer to the full pres You must refer to the full prescribing information before administering Sabril. Further information including full product data sheet is available from the Licence Holder: Marion Merrell Dow Ltd., Lakeside House, Stockley Park, Uxbridge, Middlesex. UB11 1BE.

1. Mumford JP.Br J Clin Pract 1988; 42 (Suppl 61): 7-9



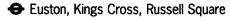
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nd includes 30% MCT for high fat absorption (87%) ensuring energy retention and enhanced protein utilizátion.

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#### References

- Committee on Nutrition of the Preterm Infant. European Society of Paediatric Gastroenterology and Nutrition. Nutrition and feeding of preterm infants. Oxford: Blackwell, 1987.

  ESPGAN Committee on Nutrition of the Preterm Infant. Acta Paediatr Scand. 1987 Suppl. 336: 1-14.

  Salle. B., Putet G. Service de Néonatologie de L'Hôpital Universitaire Edouard Herriot, Lyons, France. Senterre. J., Rigo. J., Service de Néonatologie de L'Hôpital Universitaire de Bavière, Liege, Belgium (data on file).

  Miall-Allen V.M., Whitelaw A.G.L. Dept. Paediatrics and Neonatal Medicine. Hammersmith Hospital, London (data on file).
- Brooke, O.G. et al. Archives of Disease in Childhood 1982, 57, 898-904. Lucas, A. et al. Archives of Disease in Childhood, 1984, 59, 722-730.
- Iron supplementation should therefore be considered from around 6 weeks.

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REFERENCES: 1. Vernoti S. et al, Archives of Disease in Childread, 1979, 54; 469-479, 2. Mara J.J. & Islan, A.C., Current Therapoutic Research, 1976, 20; No.1, 45-52.

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#### Alvedon™ Suppositories 125mg

PRESENTATION: White to yellowish white suppositories containing 125mg paracetamol.

**USES:** For treatment of mild to moderate pain and pyrexia in children. Alvedon suppositories may be especially useful in patients unable to take oral forms of paracetamol eg. post-operatively or with nausea

DOSAGE AND ADMINISTRATION: Children: 1-5 years, 1-2 suppositories.

The dosage should be based on age and weight ie.

- 1 year (10kg) 1 suppository 5 years (20kg) 2 suppositories

These doses may be repeated up to 4 times daily.

#### **CONTRA-INDICATIONS, WARNINGS, ETC:**

**Contra-indications:** 

Hypersensitivity to paracetamol.

**Precautions:** 

Paracetamol should be given with care to patients with impaired kidney or liver function.

Drugs which induce hepatic microsomal enzymes such as alcohol, barbiturates and other anticonvulsants may increase the hepatotoxicity of paracetamol particularly after overdosage.

Side-effects:

Side-effects at therapeutic doses are rare. Isolated cases of liver damage and allergic reactions such as skin rash have been reported.

Redness of the mucous membrane of the rectum and minor local vascular changes have been reported after the use of Alvedon suppositories. Hepatic necrosis may occur after overdosage. (See below).

#### **Pregnancy and Lactation:**

Not applicable.

PHARMACEUTICAL PRECAUTIONS: Store below 25°C.

**LEGAL CATEGORY: P** 

PACKAGE QUANTITIES: 10 strips each containing 5 suppositories or 2 strips each containing 5 suppositories.

FURTHER INFORMATION: Treatment of overdosage: Clinical symptoms of liver damage are manifested usually after 48 hours. Overdosage results in saturation of the conjugation capacity of the liver and irreversible binding of a reactive intermediate metabolite in the hepatocytes. N-acetylcysteine intravenously or L-methionine orally protects the liver if adminstered within 10-12 hours of ingesting an overdose.

PRODUCT LICENCE NO: 0017/0250.

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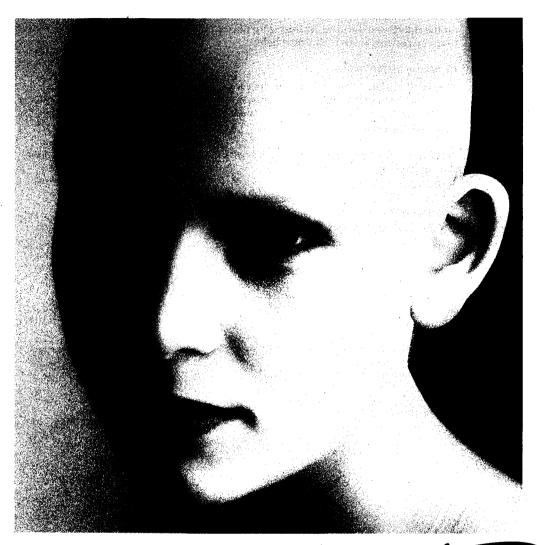
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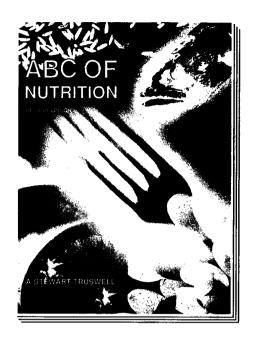


Prescribing information: Uses Nausea and vomiting due to chemotherapy or radiotherapy. Dosage Emetogenic chemotherapy and radiotherapy: Either, 8mg i.v. as a slow injection immediately before treatment, or 8mg orally 1 to 2 hours before treatment, followed by 8mg orally twelve-hourly. To protect against delayed emesis, Zofran should be continued orally, 8mg twice daily for up to 5 days. Highly emetogenic chemotherapy: A single dose of 8mg i.v. as a slow injection immediately before chemotherapy, followed by 8mg orally twice daily for up to 5 days to protect against delayed emesis. The efficacy of Zofran over the first 24 hours of highly emetogenic chemotherapy may be enhanced by the addition of a single i.v. dose of 20mg dexamethasone immediately before treatment. Alternatively, higher doses of Zofran may be beneficial, up to 32mg depending on the severity of the emetogenic challenge. Children: A single i.v. dose of 5mg/m² immediately before chemotherapy, followed by 4mg arally twelve-hourly for up to 5 days. Elderly and patients with renal impairment: No alteration of dosage, dosing frequency or route of administration is required over 65 years or with renal impairment. Patients with hepatic impairment: In patients with moderate or severe hepatic impairment, a total daily dosage of 8mg should not be exceeded. Contra-

indications Hypersensitivity to components. Precautions Pregnancy or lactation. Side effects Headache, constipation, a warm or flushing sensation in the head or epigastrium. Occasional transient rises in aminotransferases. Rare, immediate hypersensitivity reactions (see data sheet). Presentations Zofran Injection ampoules containing 4mg andansetron in 2ml aqueous solution or 8mg andansetron in 4ml aqueous solution (Product licence number 0004/0375, 4mg x 5 ampoules £52-50, 8mg x 5 ampoules £75). Zofran 4mg Tablets each containing 4mg andansetron (Product licence number 0004/0376, 4mg x 30 tablets £187-50). Zofran 8mg Tablets each containing 8mg andansetron (Product licence number 0004/0377, 8mg x 10 tablets £90). Product licence holder Glaxo Operations U.K. Limited, Greenford, Middlesex UB6 OHE. Zofran is a Glaxo trade mark. Further information is available on request from: Glaxo Laboratories Limited, Stockley Park West, Uxbridge, Middlesex UB11 IBI. Tel: 081-990 9444.



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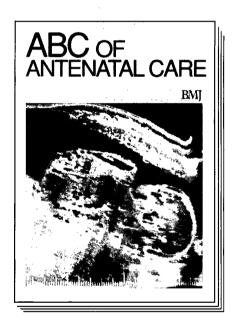
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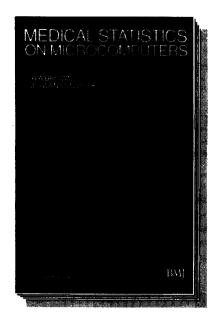
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- "... in the course of a year at least half the journals on the shelves are never looked at at all."
- "Most editors . . . are 'half nun, half whore."
- "... in the 300 years of their existence journals have done little to examine their own practices."
- "What are editors?"
- "'If an editor does not please himself,' he used to say, 'he will not please anybody.""
- "Defensive editing we do not need."
- "Medical journals in developing countries are, with very few exceptions, unattractive."
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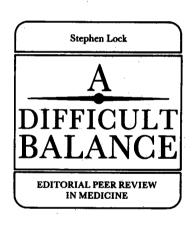
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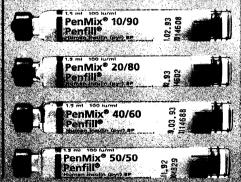
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