25000 lipase units

in just one capsule

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

467 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 proceeding.

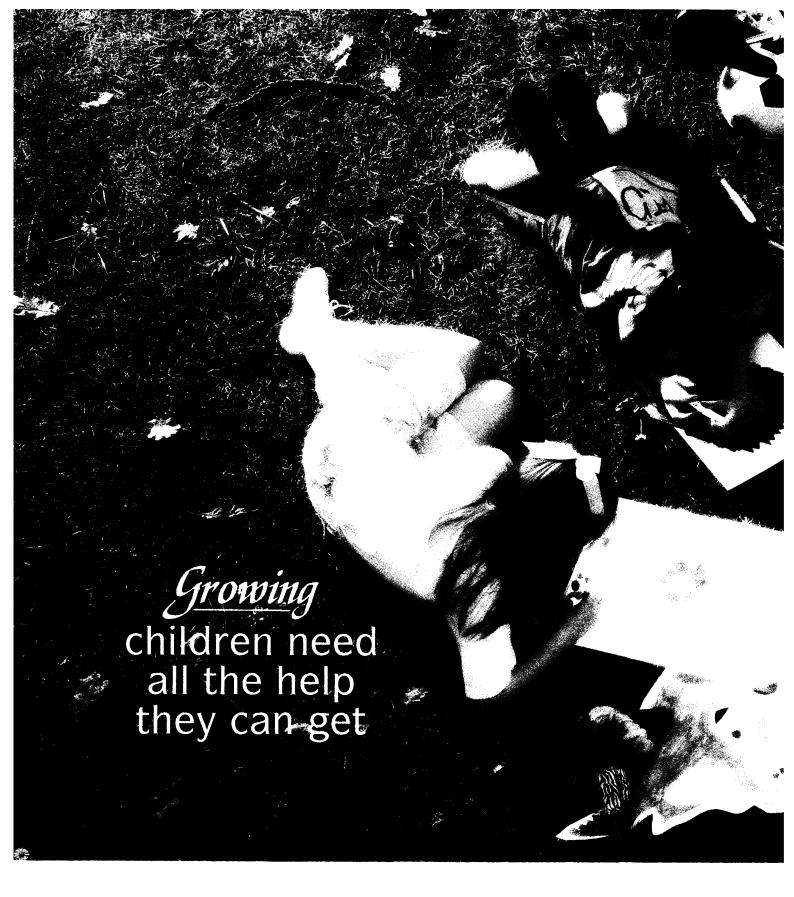
Overdosage although not experienced until now, could precipitate meconium ileus equivalent. Perianal irritation, and rarely, inflammation, could occur when large doses are used.

Product Licence Number: 5727:0006 Name and Address of Licence Holder

Kali Chemic Pharma GmbH, Hans-Bockler-Allee 20, 3000, Hannover 1, Germany. Further information is available from: Duphar Laboratories Limited, Gaters Hill, West End, Southampton SO3 3JD, Tel: (0703) 472281

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duphar



Children with growth failure need the best care you can provide. And sometimes more.

The Lilly Growth Initiative is a new support programme aimed at everyone involved in the management of growth failure. The programme looks beyond the role of growth hormone therapy, to help create the right environment for growth.

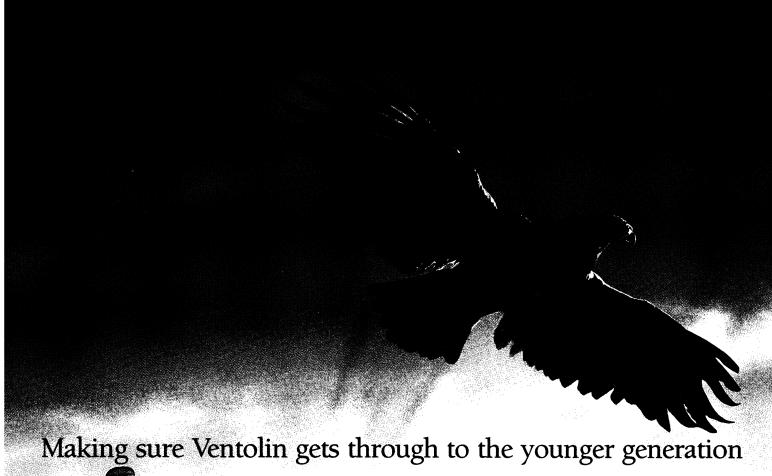
It involves communicating with children at their own level, providing advice and reassurance in terms they can easily relate to. Helping parents to understand and deal with the challenges their children face. Offering practical assistance and material support to Growth Centres. Helping clinicians through educational support, growth data management, and information services.

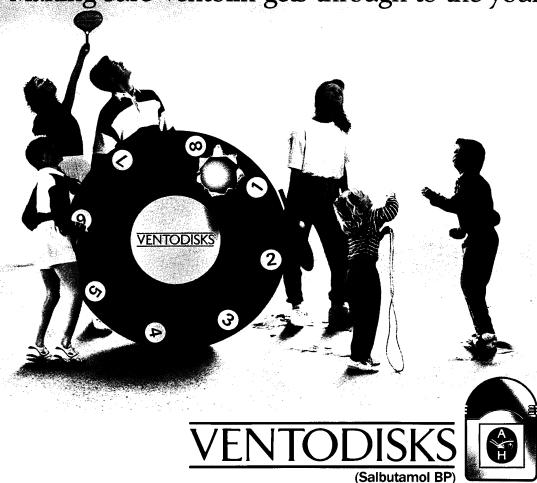


Discover how the Lilly Growth Initiative can help you succeed in the management of growth failure. Write to: The Lilly Growth Initiative, Eli Lilly & Company Ltd., Dextra Court, Chapel Hill, Basingstoke, Hampshire RG21 2SY

The Lilly Growth Initiative. A growing involvement in child development.







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. 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 β_r -agonist therapy. Paradoxical bronchospasm could occur – substitute alternative therapy. Rare reports of hyperactivity in children. **Presentation and Basic NHS cost** Hospital pack of 5 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 Ventolin Diskhaler. For inhalation – £2:54 and £4:30. **Product licence numbers** Ventodisks 200 micrograms 0045/0134, Ventodisks 400 micrograms 0045/0135.





Caring for nature's little things



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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.

Kabi Pharmacia

FIRST LINE AMINO ACIDS FOR NEONATES AND INFANTS

ABBREVIATED PRESCRIBING

L-Arginine

INFORMATION
PRESENTATION: A solution for intravenous nutrition, specifically formulated for paediatric use. I-Alanine

L-Aspartic acid	4.19
L-Cysteine/cystine	1.00
L-Glutamic acid	7.19
Glycine	2.1
L-Histidine	2.19
L-Isoleucine	3.19
L-Leucine	7.09
L-Lysine	5.69
L-Methionine	1.39
L-Phenylalanine	2.79
L-Proline	5.69
L-Serine	3.89
Taurine	0.39
LThreonine	3.69
L-Tryptophan	1.49
LTyrosine	0.59
L-Valine	3.69
In each 1000ml	65.39
Osmolality: 510mosmol	per k
water.	
Mitrogen nor litro, 03/	

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 where sufficient enteral nutrition is

impossible or impracticable
DOSAGE AND ADMINISTRATION:
Recommended dosage for infants: the dosage should be increased gradually during the first increased gradually during the first, week of administration to a final daily 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" (m)
	bodyweight i
eight (kg)	24 hours)
)	24.0
)	18.5
	16.0

40 14.5 *The duration of infusion should be

at least 8 hours **Caution:** The recommended infusion rate should not be exceeded.

Contra-indications, warnings
etc: Vaminolact is contra-indicated in patients with irreversible liver 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.

Precautions: Hyperkalaemia, hyper natraemia, and acidosis should be corrected prior to commencement of intravenous nutrition. Serum electrolytes, blood glucose levels and acid-base balance 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: Varninolact is well 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.

Package quantities: Bottles of 100ml and 500ml.

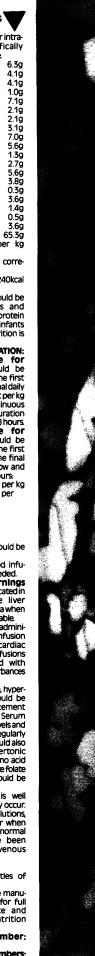
Further information: The manufacturer can be consulted for full

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

Kabi Pharmacia Limited, Davy Avenue, Kaol Pharmaca Limited, Day Avenue, Knowlhill, Militon Keynes, MK5 8PH. Distributed by Cahill May Roberts Limited for Kabi Pharmacia (ireland), Pharmapark, Chapelizod, Dublin 20. Cost: Vaminolact 100ml £3.80. Varninolact 500ml £8.70





EPILEPSY

It's not only the patient who has to put up with the disruption of epilepsy. Friends,

family, colleagues and strangers can also be

affected. Alienation and isolation can result.

For many people with epilepsy, reducing the severity of their seizures could help overcome the alienation and may be just as important as reducing the overall number of seizures.

For some patients with partial or secondarily generalised tonic-clonic seizures previously treatment-resistant. Lamictal as add-on therapy has been shown to reduce both seizure frequency and seizure severity1. Noticeably improving life for patients and those around them.

YLamictal

Making a difference to epilepsy management

Prescribing Information

Presentation: Pale yellow tablets containing 50mg and 100mg lamotrigine and coded 'LAMICTAL 50' AND 'LAMICTAL 100' respectively

Uses: Lamictal is an antiepileptic drug, indicated as an add-on treatment of partial seizures and secondarily generalised tonic-clonic seizures not satisfactorily controlled with other antiepileptic drugs. Dosage and administration: Adults, children over 12 years: Initial dose of 50mg twice a day for the first two weeks. Usual daily maintenance dose is 200-400mg in two divided doses. For patients taking sodium valproate, either alone or in combination, the initial dose is 50mg per day for the first two weeks, with a usual daily maintenance dose of 100-200mg in two divided doses. Children under 12, the elderly: As yet, insufficient information available and consequently not yet recommended for use in these groups.

Contra-indications, warnings, etc: Known hypersensitivity to lamotrigine. Cannot yet be recommended in renal/hepatic impairment. Close monitoring (including hepatic, renal and clotting parameters) of patients who acutely develop any combination of unexplained rash, fever, flu-like symptoms, drowsiness or worsening seizure control is recommended. Precautions: Liver enzyme inducers (eg. carbamazepine, phenytoin) may increase dose requirements. Sodium valproate may reduce metabolism of lamotrigine. In pregnancy the potential benefits must be weighed against any possible risk. Withdraw slowly over 2 weeks to avoid rebound seizures. Side- and adverse effects: In double-blind add-on clinical trials, skin rashes (usually maculopapular) occurred in 10% of patients taking lamotrigine and 5% of patients taking placebo. Rash led to withdrawal of lamotrigine in 2% of patients. Rarely severe skin reactions including angioedema and Stevens-Johnson syndrome have been reported. Adverse experiences normally associated with standard antiepileptics (including diplopia, dizziness, ataxia, irritability/aggression, etc.) have also been reported during trials of Lamictal when added on to such therapy. (Refer to Data Sheet for further information).

Basic NHS costs: £32.64 for pack of 56 x 50mg tablets (PL3/0273). £56.32 for pack of 56 x 100mg tablets (PL3/0274). Lamictal is a trade mark of The Wellcome Foundation Ltd.

Further information and a data sheet will be sent to any practitioner upon written request

The Wellcome Foundation Ltd., Crewe Hall, Crewe, Cheshire CW1 1UB.

Reference 1: Smith D et al. (1991). Epilepsia 32 Suppl 1: 54.







- 1. Effectiveness a 98% clinical response in otitis media or tonsillitis/pharyngitis?
- 2. Acceptability a side effect profile comparable to amoxycillin or penicillin?
- 3. Simplicity a once daily, 3 day course that is convenient for parent and child.

It is this three-sided approach that makes ZITHROMAX so suitable for children's infections.

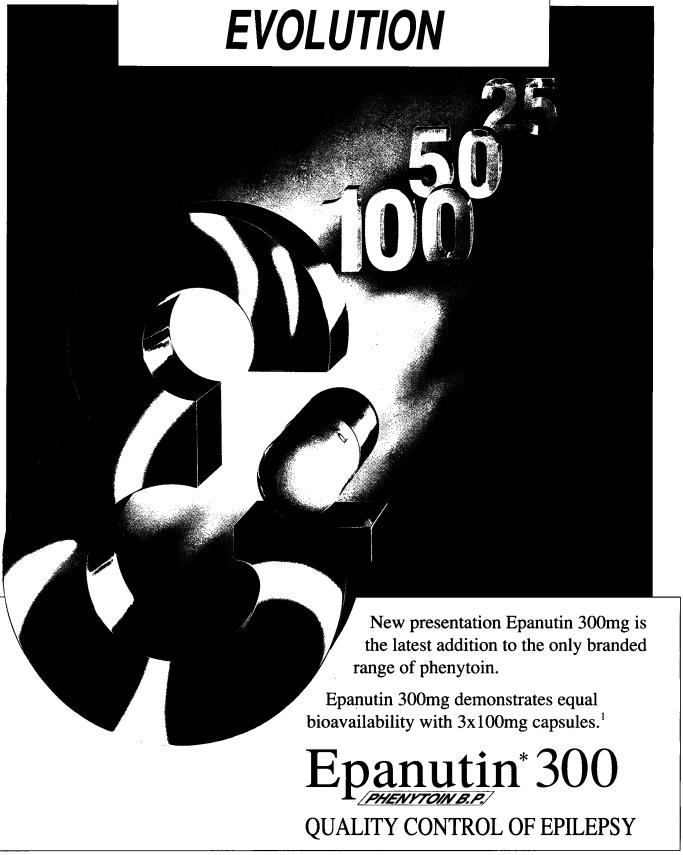
And that's why, when you treat with ZITHROMAX, you put children inside the triangle of trust.

TITHROMAX, a macrolide antibiotic, overcomes the three-sided problem of treating children's infections. 2 Ithromax.

Once daily for just 3 days

ZITHROMAX* Abbreviated Prescribing Information. Indications and dosage: Upper and lower respiratory tract infections. skin and soft tissue infections and otitis media: 500mg once daily for 3 days. Uncomplicated sexually transmitted diseases caused by Chlamydia trachomatis: Single 1g dose. Use in the elderly: Normal adult dosage is recommended. Use in children: Once daily for 3 days - Less than 3 years (up to 15kg). 10mg (0.25ml) per kg per day; 3-7 years. 5ml per day: 8-11 years. 7.5ml per day; 12-14 years. 10ml per day. There is no information on children under six months of age. See data sheet for further information on dosage recommendations according to age and weight range. Administration: ZITHRO-MAX oral suspension should be administered to children using the spoon provided in the packs or the oral dosing syringe provided in the 15ml pack only. Refer to data sheet for appropriate pack six and dispensing instructions. Contra-indications: Hypersensitivity to azithromycin or other macrolide antibiotics. Patients receiving crept derivatives. Warnings and Precautions: Moderate or severe renal impairment (creating clearance <40ml/min), liver impairment. Pregnancy and lactation: Not recommended. Drug Interactions: Antacids, ergot derivatives. Monitor patients on concurrent warfarin, digoxin or cyclosporin. Side-Effects: Nausea, abdominal discomfort, vomiting, flatulence, diarrhoea, loose stools. Allergic reactions or fash and rare reports of angioneurotic oedema and anaphylaxis: elevation in liver transaminases and reduction in neutrophil counts. Package quantities and Basic NHS Cost: 250mg capsule, pack of 6, £14.99. pack of 4, £9.99 (PL 0057/0335); powder for oral suspension - bottles of 15ml, 22.5ml and 30ml containing ZITHROMAX 200mg/5ml-15ml bottle (600mg), £5.08: 22.5ml (900mg), £7.62: 30ml (1200mg), £13.80 (PL 0057/0336). Hospital prices are available on request. In Mohs E (1992) Abstracts of the 8th Mediterranean Congress of Chemotherapy, 24th-29th May 1992: Athens, Greece. 2. Hamill J. (1992) Abstracts





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:**

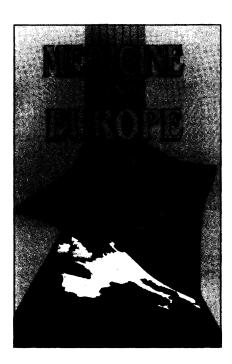
1. Data on file, Parke-Davis Research Laboratories.

PARKE-DAVIS TESTICAL

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THE WAY AHEAD



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Published June 1992

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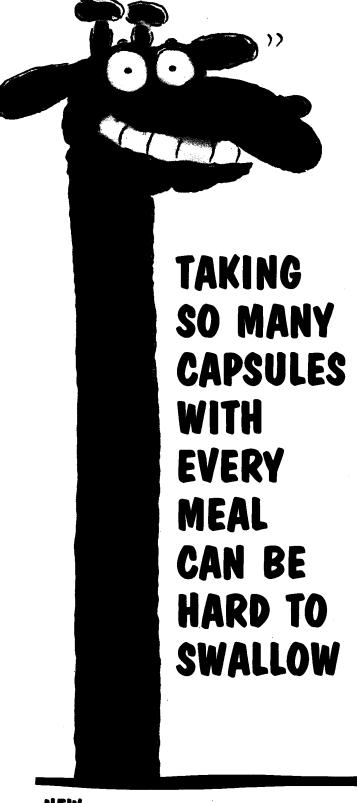
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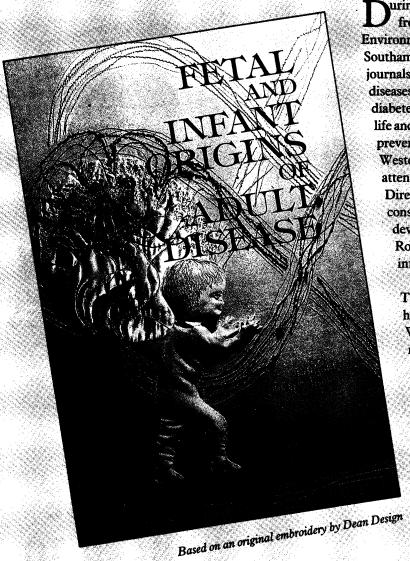


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uring the past few years a series of articles, mostly from the Medical Research Council's Environmental Epidemiology Unit at the University of Southampton, has been published in leading medical journals. They set out the evidence that certain adult diseases, including coronary heart disease, stroke and diabetes originate in impaired development during fetal life and infancy. Because of the obvious implications for prevention of some of the commonest diseases in Western society, they have attracted international attention. In this book, Professor David Barker, Director of the Unit, has selected 31 articles that he considers seminal and a comprehensive guide to the development of this important topic. Professor Roger Robinson's introduction summarises and interprets the evidence for non-epidemiologists.

> The first chapters describe the origins of the hypothesis in geographical studies in England and Wales. These are followed by a series of studies of men and women in middle and late life whose early growth was recorded at the time. In those who have died, cause of death can be related to early growth. Examination of the living has allowed blood pressure, blood lipid and insulin concentrations, and other measurements to be related to different patterns of early growth. Together, the findings show that early development affects the risk of coronary heart disease, stroke, obstructive lung disease and diabetes at least as strongly as obesity, smoking and other aspects of adult life style.

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Sachets: Containing 500 mg vigabatrin powder and no other ingredients. Dissolve in water or soft drink immediately before use. Uses: Indications: Treatment of epilepsy not controlled by other antiepileptic drugs. Dosage and 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 between plasma concentration and efficacy. 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 30-50kg 1.5-3g/day 30-50kg 2-4g/day

Date of Preparation: September 1992.

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.

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



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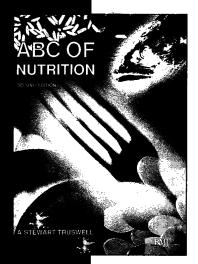
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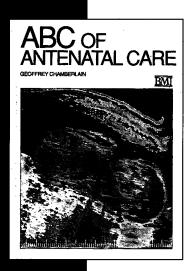
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American Journal of Clinical Nutrition

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January 1992

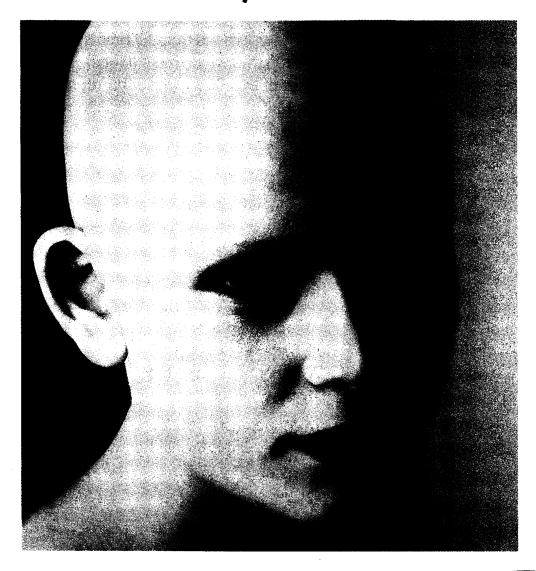
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Yet, there is no evidence of extra-pyramidal reactions, 2,3,4 little risk of unwanted sedation3,4 and no reported drug interactions.4

At a time when patients need all the help they can get, Zofran will help make the treatment easier to bear.



Prescribing Information: Uses: Nausea and vomiting due to chemotherapy or radiotherapy, and postoperative nausea and vomiting. **Dosage:** Emetogenic chemotherapy and cadiotherapy: Either, 8mg i.v. as a slow injection immediately before treatment, gr 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 emetagenic 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 orally twelve-hourly for up to 5 days. *Elderly and patients with renal*

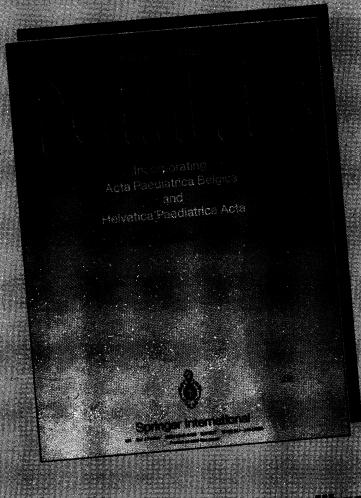
impairment: No alteration of dosage, dosing frequency or route of administration is required over 65 years or with renal impairment. Patients with hepatic impairment: In patients with moderate or severe hepatic impairment, a total daily dosage of 8mg should not be exceeded. Postoperative nausea and vomiting: For prevention in adults: Either a single dose of 4mg i.v. as slow injection at induction, or 8mg orally I hour prior to anaesthesia followed by two further 8mg doses at eight-hourly intervals. For treatment in adults: A single dose of 4mg i.v. as a slow injection. There is no experience in postoperative nausea and vomiting in children and limited experience in the elderly. The role of ondansetron in opiate-induced emesis is not yet established. 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 Amg ondansetron

in 2ml aqueous solution or 8mg ondansetron in 4ml aqueous solution (Product licence number 0004/0375, 4mg x 5 ampoules \$52-55; 8mg x 5 ampoules \$75). Zofran 4mg Tablets each containing 4mg ondansetron (Product licence number 0004/0376, 4mg x 30 tablets £187-50), Zofran 8mg Tablets each containing 8mg ondansetron (Product licence number 0004/0377, 8mg x 10 tablets £90). Product licence holder: Glaxo Operations UK 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 UBI1 1BT. Tel: 081-990 9444.

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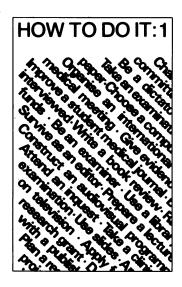
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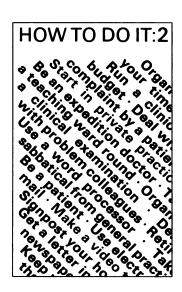
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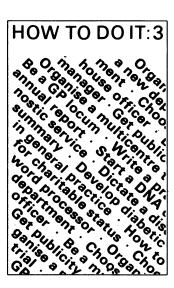


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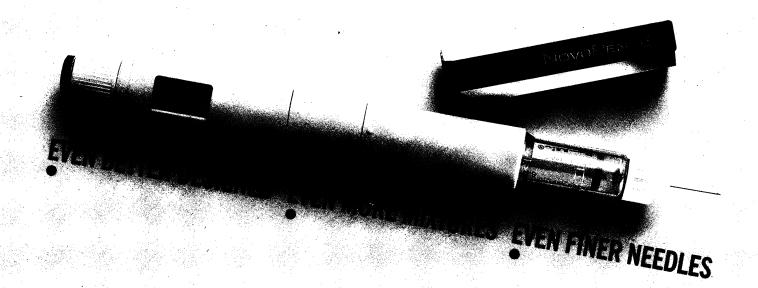
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