

# A NEW PRESENTATION

# ATROVENT

ipratropium bromide

# Autohaler<sup>®</sup>

Add a boost to your  
bronchodilator therapy



#### PRESCRIBING INFORMATION

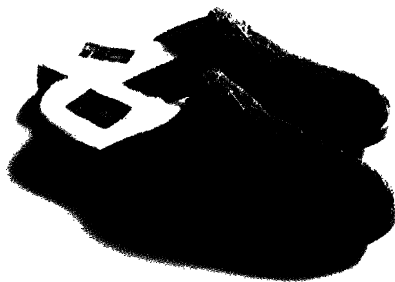
**ATROVENT AUTOHALER** Ipratropium bromide. Indications Chronic reversible airways obstruction, particularly chronic bronchitis. Dosage *Adults*: Up to 4 puffs three or four times daily. *Children 6-12 years*: 1 or 2 puffs three times daily. *Under 6 years*: 1 puff three times daily. **Contra-indication** Known hypersensitivity to atropine. **Precautions** Glaucoma; prostatic hypertrophy; pregnancy, especially the first trimester. Advise patients to seek medical advice if response lessens. **Side effects** Dry mouth may occur. **Presentation** Breath-actuated pressurised aerosol for inhalation therapy. 10ml vial complete with mouthpiece contains 200 doses, each delivering 20 micrograms ipratropium bromide £10.43. **Legal category** POM. PL 0015 0160. **Product licence holder** Boehringer Ingelheim Ltd, Ellesfield Avenue, Bracknell, Berkshire RG12 8YS. For full prescribing information please see data sheet. Date of preparation September 1993.



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Ingelheim**

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Oral Ciproxin® brings you parenteral power in a tablet, a simple choice that can save hospital time and expense in the treatment of a range of infections.

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As it promotes rapid recovery, oral Ciproxin® helps get the patient back on his feet – fast.

**ABRIDGED PRESCRIBING INFORMATION CIPROXIN® INFUSION/TABLETS** (Refer to data sheet before prescribing). **Presentation Infusion:** Clear, almost colourless solution containing the equivalent of 100 mg ciprofloxacin in 50 ml, and 200 mg ciprofloxacin in 100 ml solution. **Tablets:** White tablets containing the equivalent of either 250 mg, 500 mg or 750 mg ciprofloxacin. **Uses** Ciprofloxacin is indicated for the treatment of single or mixed infections caused by susceptible organisms. Also indicated for prophylaxis against infection in elective upper gastro-intestinal surgery and endoscopy where there is an increased risk of infection. **Dosage and administration Adults** Ciproxin Infusion 100–200 mg administered intravenously over 30 to 60 minutes twice daily. Ciproxin Tablets should be swallowed whole with liquid; the dosage range for adults is 250–750 mg twice daily. In surgical prophylaxis a single 750 mg tablet administered 60–90 minutes before the procedure (but see Interactions with oral premedicants). **Elderly** No dose adjustment. **Duration of treatment** For acute infections the usual treatment period is 5 to 7 days (intravenous) or 5 to 10 days (oral), except in cases of acute uncomplicated cystitis where treatment is 250 mg twice daily for 3 days (oral). Generally, in acute and chronic infections where sensitivity is proven, treatment should be continued for at least 3 days after the signs and symptoms of infection have disappeared.

**Contra-indications** Hypersensitivity to ciprofloxacin or other quinolones; also in children and growing adolescents except where the benefits of treatment outweigh the risks. **Warnings and precautions** Use with caution in epileptics and patients with a history of CNS disorders. Treatment could result in impairment of the ability to drive or operate machinery. Crystalluria has been reported so patients should be well hydrated and excessive urine alkalinity avoided. As haemolytic reactions with ciprofloxacin are possible in patients with latent and actual defects in glucose-6-phosphate dehydrogenase activity, use with caution. **Drug interactions** Increased plasma levels of theophylline have been



s t w a y

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t o



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**oral** **Ciproxin**<sup>®</sup>  
ciprofloxacin

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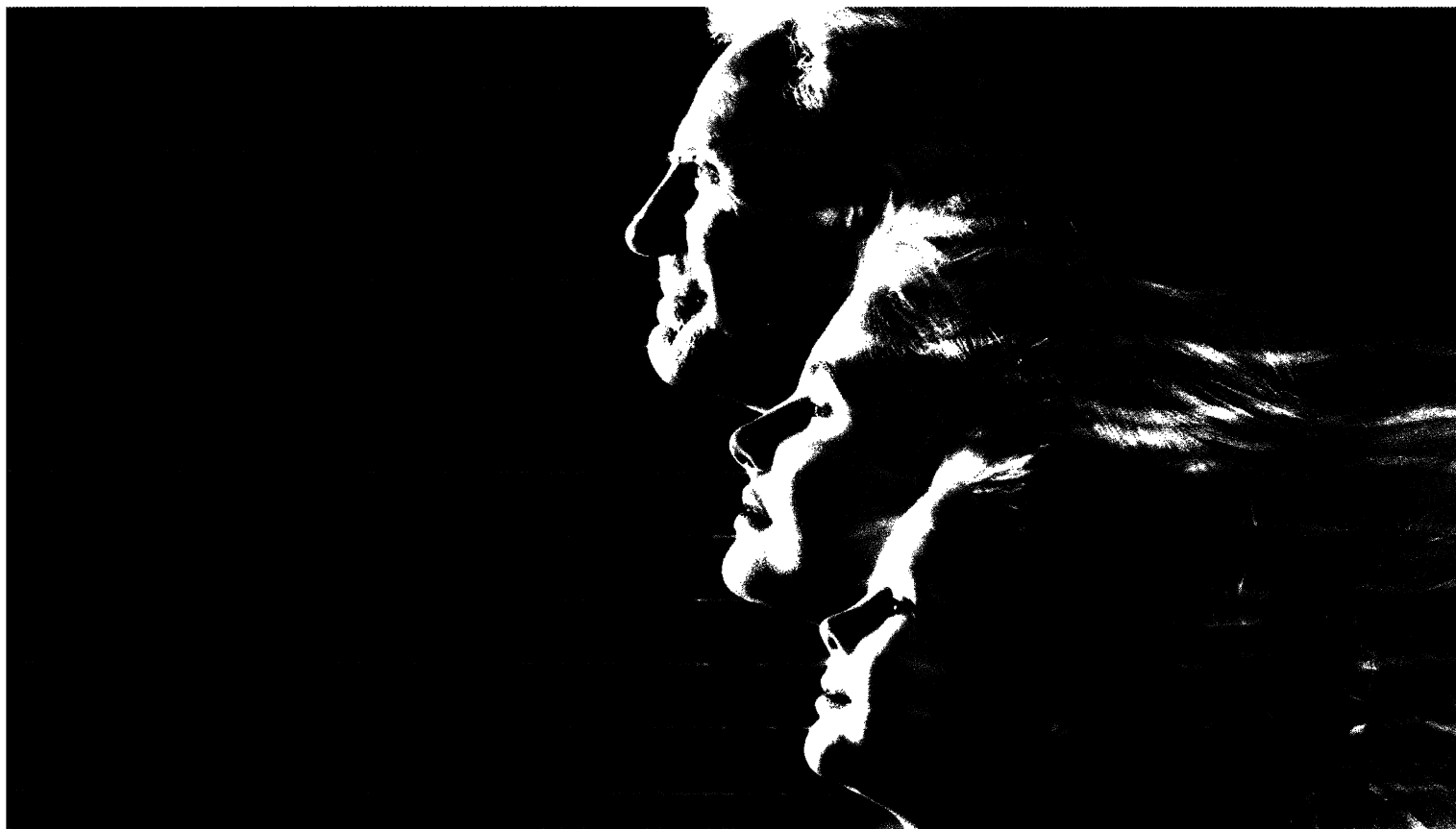
**Parenteral power in a tablet.**

observed following concurrent administration with ciprofloxacin. The dose of theophylline should be reduced and plasma levels of theophylline monitored. Where monitoring of plasma levels is not possible, avoid the use of ciprofloxacin in patients receiving theophylline. Particular caution is advised in those patients with convulsive disorders. Interactions have also been noted with anticoagulants and cyclosporin. High doses of quinolones have shown an interaction with NSAIDs in animals leading to convulsions. Administration of quinolones and glibenclamide simultaneously can potentiate the effect of glibenclamide, resulting in hypoglycaemia. Opiate premedicants or regional anaesthetic agents must not be administered concomitantly with ciprofloxacin when used for surgical prophylaxis. Ciproxin Tablets should not be administered within 4 hours of medications containing magnesium, aluminium or iron salts. **Use in pregnancy and lactation** Not recommended. **Side-effects** Gastro-intestinal, CNS, hypersensitivity/skin reactions, musculoskeletal and special sense disturbances. Renal and hepatic disturbances. Effects on haematological parameters. Also reported: vasculitis, pseudomembranous colitis, Stevens-Johnson Syndrome, Lyell Syndrome, haemolytic anaemia, granulocytopenia, intracranial hypertension, petechiae, haemorrhagic bullae, tenosynovitis and tachycardia. Local irritation at the site of injection (infusion only). **Overdosage** Serum levels of ciprofloxacin are reduced by dialysis. **Pharmaceutical precautions** Unless compatibility is proven, the infusion should always be administered separately. Do not refrigerate Ciproxin Infusion. **Legal category** POM. **Package quantities** Ciproxin Infusion bottles of 50 or 100 ml. Ciproxin Tablets Blister strips of 10 in packs of 10, 20, and 100. **Product licence numbers** PL 0010/0150 Infusion. PL 0010/0146-0148 Tablets. **Basic NHS cost** 250 mg x 10 tablets £ 7.50. 500 mg x 10 tablets £ 13.75. 750 mg x 10 tablets £ 20.00. 100 mg infusion £ 12.00. 200 mg infusion £ 24.00. **Date of preparation** July 1993. Baypharm, Bayer plc, Pharmaceutical Division, Bayer House, Strawberry Hill, Newbury, Berkshire RG13 1JA, Tel: (0635) 39000. © Registered trademark of Bayer AG, Germany.

**Presentations:** *Pulmicort Respules* (2 ml single dose unit ampoules) containing 0.25 mg/ml or 0.5 mg/ml budesonide in a suspension for nebulisation. **Uses:** Bronchial asthma where use of a pressurised inhaler or dry powder formulation is unsatisfactory or inappropriate. **Dosage and administration:** Dosage schedules: Administer from suitable nebulisers. Dose delivered to the patient varies depending on the nebulising equipment used (see data sheet). Adjust dosage individually. Initially during periods of severe asthma and while reducing or discontinuing oral glucocorticosteroids the recommended dose in adults (including elderly and children 12 years and older) is usually 1-2 mg twice daily. In very severe cases the dosage may be further increased. Children 3 months to 12 years: 0.5-1 mg twice daily. The maintenance dose should be the lowest dose which keeps the patient symptom-free. Recommended doses are: Adults (including elderly and children 12 years and older): 0.5-1 mg twice daily. Children (3 months to 12 years): 0.25-0.5 mg twice daily. For an increased therapeutic effect increase dose of *Pulmicort* rather than combine treatment with oral corticosteroids because of the lower risk of systemic effects. **Contra-indication:** Hypersensitivity to any of the constituents. **Special warnings and precautions:** Care is needed in patients with pulmonary tuberculosis and viral infections in the airways. A short course of oral steroids in addition to *Pulmicort* may be required in patients with excessive mucus in



the bronchi. Transfer of patients dependent on oral steroids to *Pulmicort* demands special care; see data sheet for further details. The nebuliser chamber should be cleaned and dried after every administration. *Pulmicort* does not affect the ability to drive and use machines. *Pulmicort Respules* can be mixed with 0.9% saline and with solutions of terbutaline, salbutamol, sodium cromoglycate or ipratropium bromide. **Side effects:** Mild irritation in the throat, coughing and hoarseness and oral candidiasis have been reported. In rare cases inhaled drugs may provoke bronchoconstriction in hyperreactive patients. Facial skin should be washed after use of the face mask as irritation can occur. Coughing can usually be prevented by inhaling a B<sub>2</sub> agonist (e.g. terbutaline) 5-10 minutes before inhalation of *Pulmicort Respules*. Avoid in pregnancy. **Pharmaceutical precautions:** Store below 30°C. Use within 3 months of opening the foil envelope. Protect opened ampoule from light. Use within 12 hours of opening. **Legal category:** POM. **Basic NHS price:** *Pulmicort Respules* 0.25 mg/ml (20 single dose units) £32.00. *Pulmicort Respules* 0.5 mg/ml (20 single dose units) £44.64. **Product licence numbers:** *Pulmicort Respules* 0.25 mg/ml PL 0017/0309. *Pulmicort Respules* 0.5 mg/ml PL 0017/0310. **For further information contact the product licence holder:** Astra Pharmaceuticals Ltd., Home Park, Kings Langley, Herts WD4 8DH. **Reference:** 1. BOSS Study, *Thorax* 1993; **48(4)**.



# Time to take a breather from oral steroids

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***Respules***<sup>®</sup>

A high-dose nebulised steroid that's low on side effects<sup>1†</sup>

# FLIXOTIDE<sup>▼</sup>

fluticasone propionate

Flixotide (fluticasone propionate)  
Abridged Prescribing Information  
(Please refer to the full data sheet  
before prescribing)

**Uses** Topically active corticosteroid  
for prophylactic management of  
asthma. **Dosage and administration**  
For inhalation only. Use regularly.  
Onset of therapeutic effect usually  
occurs in 4 to 7 days. **Adults:** 100 to  
1,000 micrograms twice daily.  
**Children over 4 years:** 50 to 100  
micrograms twice daily.

**Contra-indication** Hypersensitivity.  
**Precautions** *Severe or unstable  
asthma:* Warn patients to seek  
medical advice if short-acting inhaled  
bronchodilator use increases or  
becomes less effective. Consider  
using oral steroids and/or maximum  
doses of inhaled corticosteroids.  
Treat severe exacerbations in the  
normal way. *Acute symptoms:*

Flixotide is not for relief of acute  
symptoms. A short-acting inhaled  
bronchodilator is required. **Systemic  
effects:** Adrenal function and reserve  
usually remain within the normal  
range. Some systemic effects may  
occur in a small proportion of adults  
after long-term treatment at  
maximum recommended dose.

No systemic side effects have been  
seen in children. **Transfer from oral  
steroids:** Special care is needed.

Monitor adrenal function. Do not  
stop Flixotide abruptly. Consider  
additional corticosteroid therapy in  
situations likely to produce stress.

**Tuberculosis:** Special care is needed  
in active or quiescent pulmonary  
tuberculosis. **Pregnancy and lactation:**  
Experience is limited. Balance risks  
against benefits. **Side effects**

Candidiasis of mouth and throat,  
hoarseness. **Paradoxical  
bronchospasm:** Substitute alternative  
therapy. **Presentation and Basic**

**NHS cost Flixotide Inhaler:** 120  
actuations per inhaler. 25 micrograms

– £6.86. 50 micrograms – £11.43. 125  
micrograms – £22.86. 250 micro-

grams – £38.86. **Flixotide Diskhaler:**

Pack of 14 four-place disks together  
with a Flixotide Diskhaler. 50 micro-

grams – £8.23. 100 micrograms

– £12.80. 250 micrograms – £24.23.

500 micrograms – £40.23. **Flixotide**

**Diskhaler refill pack:** Pack of 14 four-  
place disks only. 50 micrograms –

£7.66. 100 micrograms – £12.23.

250 micrograms – £23.66. 500  
micrograms – £39.66. Hospital packs

are also available. **Product licence**

**numbers** 10949/0001, 10949/0002,  
10949/0003, 10949/0004,

10949/0005, 10949/0006,  
10949/0007, 10949/0008.

**References** 1. Phillipps GH.  
Structure-activity relationships of  
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ALLEN & HANBURY'S

Further information is available  
on request from:

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## PUTTING A SMILE ON THE FACE OF ASTHMA

Flixotide, a significant new inhaled steroid from Allen & Hanburys,  
has been developed to help put a smile on the face of asthma.

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inflammatory activity<sup>1</sup> with negligible oral systemic bioavailability.<sup>2</sup>  
Flixotide is effective where it's needed in the lungs with minimal potential  
for steroid side effects from the swallowed portion.



#### Abridged Prescribing Information.

**Presentation:** *Rhinocort Aqua*: A metered pump spray delivering 100 µg budesonide per dose. *Rhinocort Nasal Aerosol*: A metered dose aerosol delivering 50 µg budesonide per dose. **Uses:** Seasonal and perennial allergic rhinitis and vasomotor rhinitis. **Dosage:** *Adults (including elderly)*: 400 µg once daily in the morning, or 200 µg twice daily, morning and evening. When good effect has been achieved, reduce dose. *Children:* *Rhinocort Aqua*: Not recommended. *Children over 6 years*: use *Rhinocort Nasal Aerosol*, dosage as for adults. **Contra-indications, warnings etc.:** Hypersensitivity to any of the ingredients. Special care demanded when treating patients transferred from oral steroids, where disturbances of hypothalamic-pituitary-adrenal (HPA) axis could be expected. Special care needed in patients with fungal and viral infections in the airways, or with lung tuberculosis. Full effect not achieved until after a few days' treatment. Treatment of seasonal rhinitis should start, if possible, before exposure to the allergens. Concomitant treatment may sometimes be necessary to counteract eye symptoms. In continuous, long-term treatment, the nasal mucosa should be inspected regularly. Continuous, long-term treatment of children is not recommended. *Rhinocort* does not affect ability to drive and operate machinery. Avoid during pregnancy. **Side-effects:** Sneezing, nasal stinging and dryness may follow immediately after use of spray. Slight haemorrhagic secretion may occur. Contact allergy involving facial skin may occur rarely. Rare cases of cataract after prolonged use have been reported. Ulceration of mucous membrane and nasal septal perforation have been reported rarely. **Package quantities and NHS cost:** *Rhinocort Aqua* — 100 x 100 µg doses budesonide INN — £6.00. *Rhinocort Nasal Aerosol* — 200 x 50 µg doses — £5.66. **Product licence No.:** 0017/0304 — *Rhinocort Aqua*. 0017/0204 — *Rhinocort Nasal Aerosol*. **Legal category:** POM. **Further information is available from:** Astra Pharmaceuticals Ltd., Home Park, Kings Langley, Herts WD4 8DH.

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Astra Pharmaceuticals Ltd.,  
Home Park, Kings Langley, Herts WD4 8DH.  
Telephone: 0923 266191.

RHA 803/04/93

Date of preparation: April 1993



- The effectiveness of budesonide has been demonstrated in both seasonal<sup>1-10</sup> and perennial<sup>11-</sup>

- 74% of patients have stated

*Rhinocort Aqua* once

\* Efficacy demonstrated in references 1 to 18 inclusive.



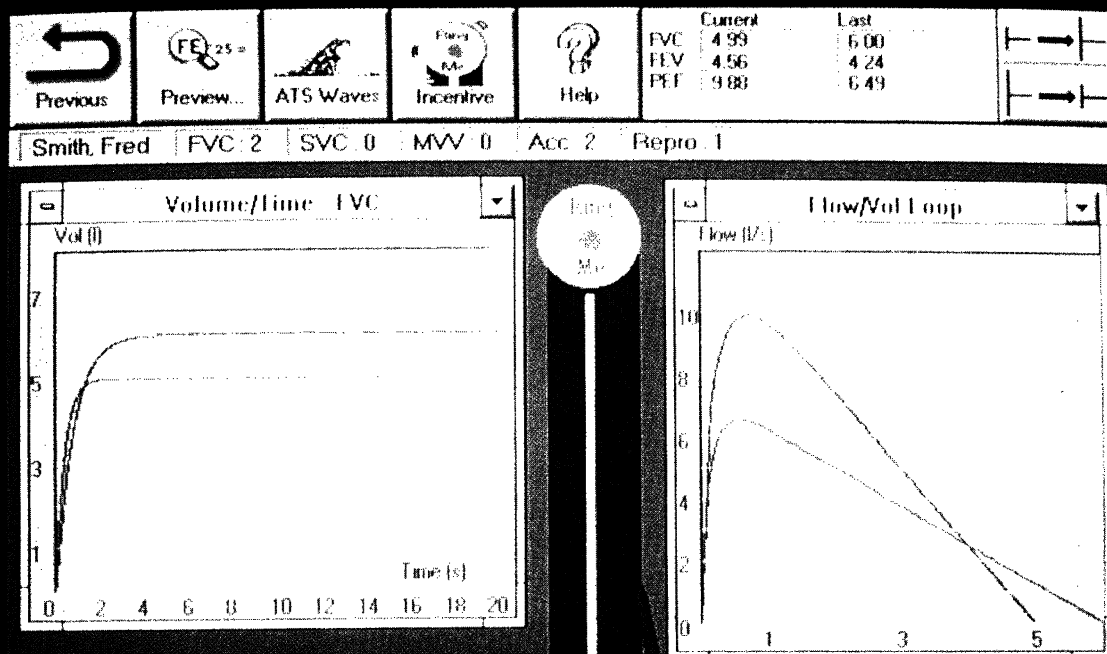
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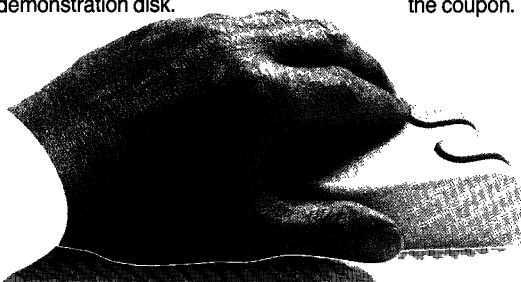
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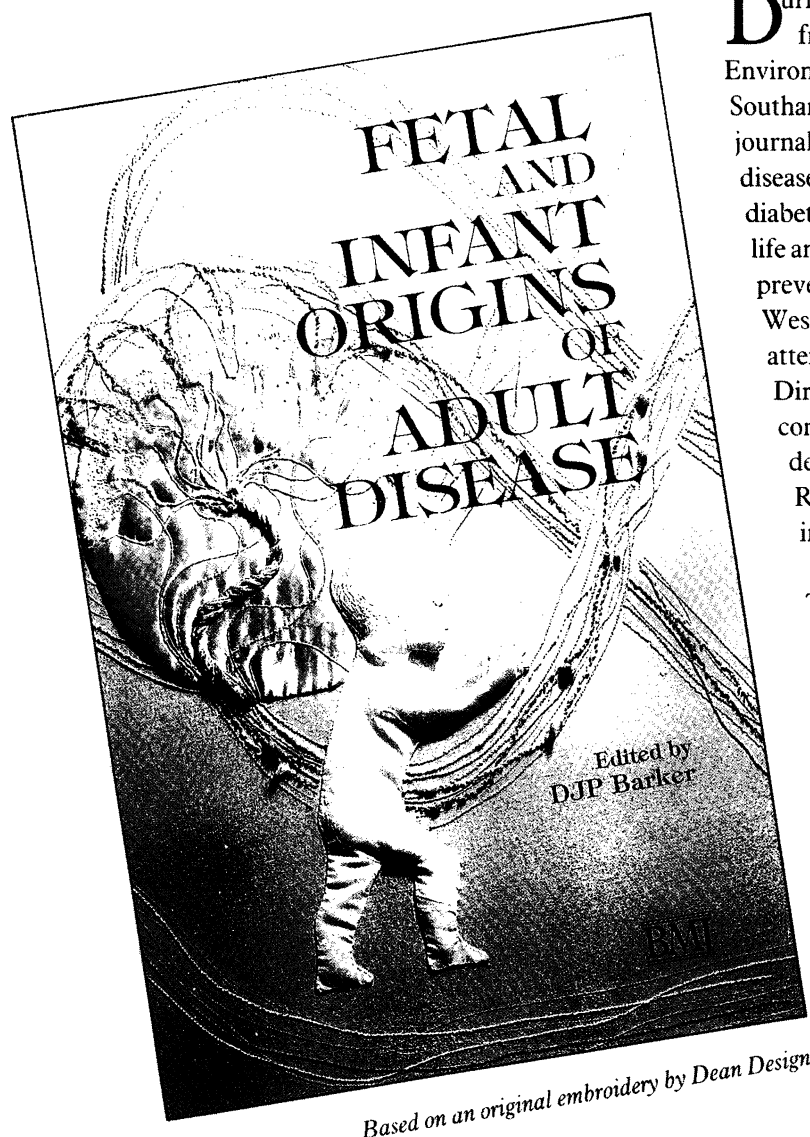
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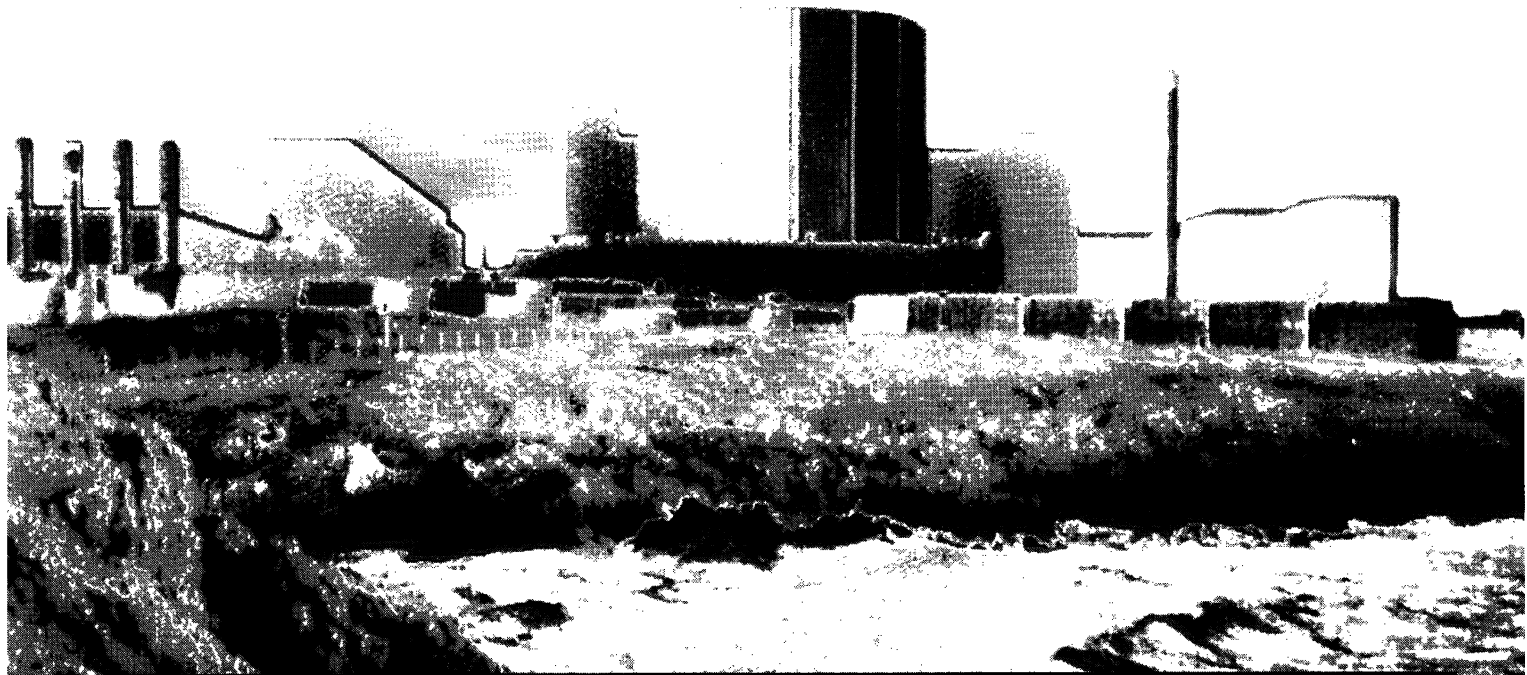
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Edited by **Valerie Beral, Eve Roman & Martin Bobrow**



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*As a result of the Gardner paper, considerable attention has been paid to the problem of childhood cancer near nuclear installations.*

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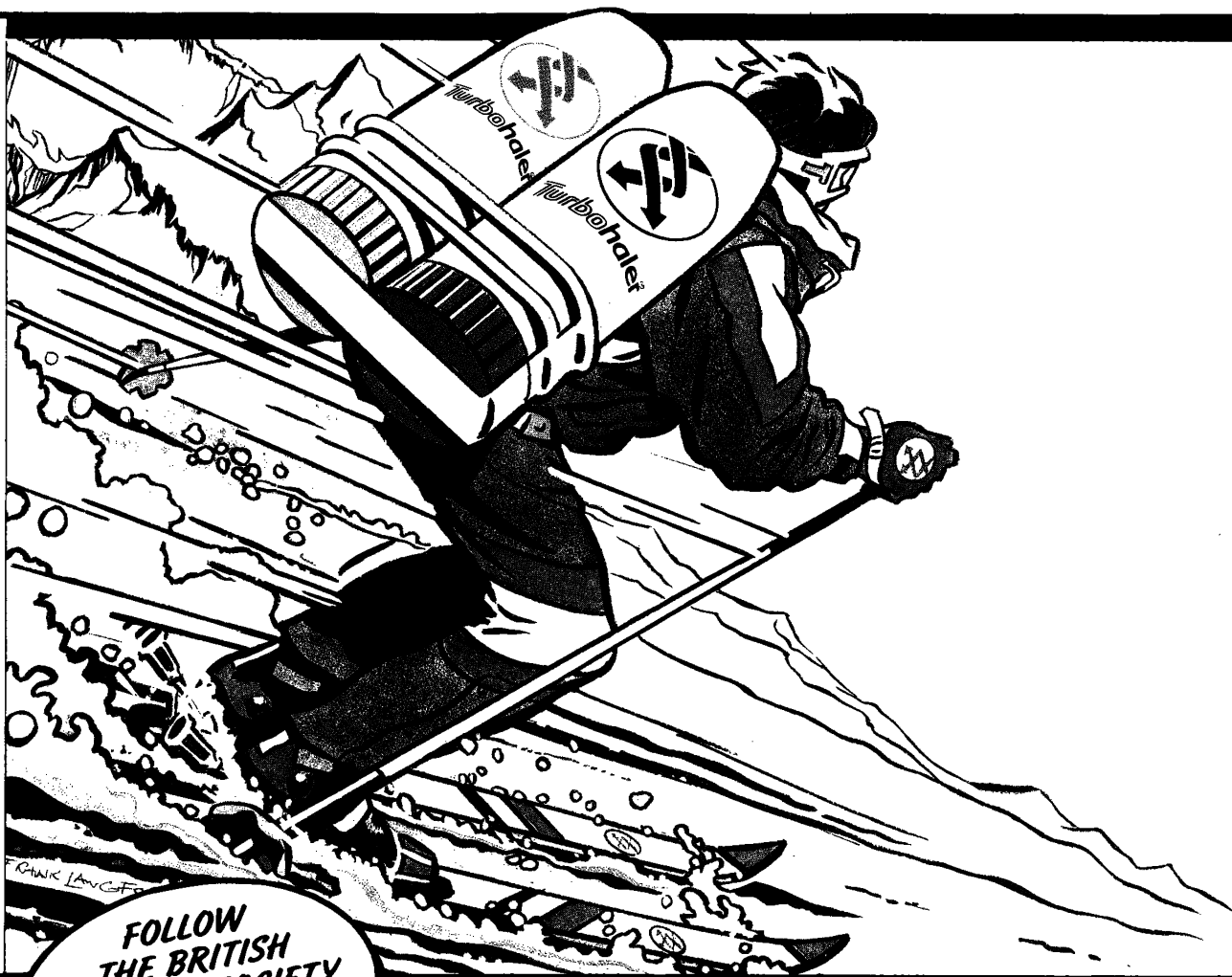
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# TURBOHALED!



**FOLLOW  
THE BRITISH  
THORACIC SOCIETY  
GUIDELINES\* FOR  
ASTHMATICS**

**STEP 1: INHALED BRONCHODILATOR PRN  
STEP 2: ADD IN LOW-DOSE INHALED STEROID**

**Bricanyl** **Pulmicort**  
 **Turbohaler**  **Turbohaler 200**  
 TERBUTALINE SULPHATE BUDESONIDE

**Abridged Prescribing Information: Presentation:** Bricanyl Turbohaler. Breath actuated metered dose powder inhaler delivering 500µg terbutaline sulphate per dose. Each inhaler contains 100 doses. **Uses:** Relief of bronchospasm. **Dosage and Administration:** Adults (including elderly) and children: One inhalation as required, up to four times daily. **Contra-indications, warnings, etc.:** Do not use in patients hypersensitive to terbutaline or with hypertrophic cardiomyopathy. Care advised in myocardial insufficiency, thyrotoxicosis and during the first trimester of pregnancy. Potentially serious hypokalaemia may result from  $\beta_2$ -agonist therapy. Caution advised in severe asthma as effect may be potentiated by concomitant treatment with xanthines, steroids, diuretics and by hypoxia (see data sheet). Do not administer with  $\beta$ -blockers and use with caution with other sympathomimetics. Additional blood glucose measurements are recommended initially in diabetic patients. Patients should be warned to seek medical advice if the usual relief or duration of action is diminished. **Side-effects:** Infrequent: tremor, tonic cramp, tension and palpitations. **Legal Category:** POM. **Basic NHS price:** Bricanyl Turbohaler, (100 doses) £8.94. **Product Licence Number:** PL 0017/0241. **For further information contact the product licence holder:** Astra Pharmaceuticals Ltd., Home Park, Kings Langley, Herts WD4 8DH.

**Abridged Prescribing Information: Presentations:** Pulmicort Turbohaler 200 – 200µg/puff dry powder inhaler containing 100 doses of

budesonide. Pulmicort Turbohaler 400 – 400µg/puff dry powder inhaler containing 50 doses of budesonide. **Uses:** Bronchial asthma. **Dosage and Administration:** Individualise dose. **Adults:** 200µg-1600µg daily in divided doses. **Children:** 200µg-800µg daily in divided doses. **Maintenance:** Use lowest possible dose. Rinse mouth after each use. **Contra-indications:** None known. **Warnings, etc.:** Active lung tuberculosis. Care is needed in patients with fungal and viral infections in the airways. Avoid administration during pregnancy. A short course of oral steroids in addition to Pulmicort may be required in patients with excessive mucus in the bronchi. Transfer of patients dependent on oral steroids to treatment with Pulmicort demands special care. See data sheet for further details. **Side-effects:** Mild irritation in the throat; hoarseness and oral candidiasis occur in some patients. Rare cases of cataract have been reported after prolonged use of corticosteroids. **Legal Category:** POM. **Licence No:** PL 0017/0271 (400µg/puff). PL 0017/0272 (200µg/puff). **Price:** Pulmicort Turbohaler 200 and Pulmicort Turbohaler 400 £18.50. **For further information contact the product licence holder:** Astra Pharmaceuticals Ltd., Home Park, Kings Langley, Herts WD4 8DH. **Reference:** 1. Statement by the British Thoracic Society, Research Unit of The Royal College of Physicians of London, King's Fund Centre, National Asthma Campaign. *Brit Med J* 1991; 301: 651-653.

**ASTRA**

Astra Pharmaceuticals Ltd., Home Park, Kings Langley, Herts, WD4 8DH

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