

A Logical Combination in **COPD**

Salbutamol + ipratropium bromide in a single metered dose inhaler

Prescribing information Combivent Metered Aerosol containing 200 doses, each delivering ipratropium bromide (anticholinergic bronchodilator) 20 micrograms and salbutamol (β)-adrenergic agonist) 100 micrograms. Indication: treatment of bronchospasm associated with chronic obstructive pulmonary disease in patients who require regular treatment with both ipratropium and salbutamol. Dosage: Adults only: two puffs four times a day. Contra-indication: known hypersensitivity to any of the components or to atropine or its derivatives. Precautions: cardiac disorders: hyperthyroidism; diabetes mellitus; co-prescription with β-blockers,

Boehringer Ingelheim

hyperthyroidism; diabetes mellitus; co-prescription with β-blockers, corticosteroids, xanthine derivatives, other β-agonists or anticholinergics; pregnancy, especially the first trimester, and breast feeding. Potentially

serious hypokalaemia may result from β:-agonist therapy. Advise patient to seek medical advice in the event of acute, rapidly worsening dyspnoea or if response lessens; do not spray into the eye. **Side-effects**: tremor and nervousness may occur; tachycardia, dizziness, palpitations, headache, local reactions such as dryness of the mouth are less frequent; urinary retention has been reported rarely. As with other bronchodilators, cough and, very rarely, paradoxical bronchoconstriction have been observed. **Basic NHS price**: 10ml vial complete with mouthpiece UK £6.00. POM. PL 0015.0191. PA 7 52·1 **Product Licence and Authorisation Holder**: Boehringer Ingelheim Limited, Ellesfield Avenue, Bracknell, RG12 8YS. For full prescribing information please contact sheet.

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Beclazone Easi-Breathe Inhaler

Beclamethasone Dipropionate BP

(Please refer to full data sheet before prescribing)

- Uses Provides automatic actuation of inhaler with inspiration. For the management bronchial asthma especially in patients inadequately controlled by bronchodilators and sodium cromoglycate.
- Dosage and Administration Use regularly. Adults, Beclazone 50 and 100 Easi-Breathe Inhalers; 100 microgram three or four times daily. Beclazone 250 Easi-Breathe Inhaler; 500 microgra twice a day or 250 microgram four times a day. Elderly, no dose adjustment necessary, including patients with renal or hepatic impairment. Children, Beclazone 50 and 100 Easi-Breathe Inhalers; 50 to 100 microgram two to four times daily. Beclazone 250 Easi-Breathe Inhaler is not indicated for use in children.
- Contra-Indications Hypersensitivity to the ingredie
- Precautions Patients should be instructed in the correct use of inhalers. May induce systemic cortico-steroid effects (with reduction in plasma cortisol levels) and adrenal suppression (above 2000 microgram daily) - monitor adrenal function and provide systemic steroids in appropriate cases of stress. Caution in patients with history of, or active pulmonary tuberculosis. Avoid sudden cessation of treatment.
- Pregnancy/Lactation Use inhalers only if the potential benefit outweighs the risk.
- Side Effects Paradoxical bronchospasm discontinue use immediately and seek medical advice. Candidiasis, hoarseness or throat irritation - relieve by rinsing throat with water.
- Presentations and Basic NHS Cost Metered-Dose Aerosol supplied in a Breath-Operated Inhaler containing 200 doses. Beclazone 50 Easi-Breathe, Beclazone 100 Easi-Breathe and Beclazone 250 Easi-Breathe Inhalers deliver 50, 100 and 250 microgram beclomethasone dipropionate per actuation of the valve.
- Product Licence Numbers (Cost) Beclazone 50 Easi-Breathe Inhaler PL 0530/0451 (£4.34) Beclazone 100 Easi-Breathe Inhaler PL 0530/0452 (£8 24) Beclazone 250 Easi-Breathe Inhaler PL 0530/0453 (£18.02)
- Legal Category POM
- Further Information is available on request from: Baker Norton Gemini House, Flex Meadow, Harlow Essex CM19 5TI
- Date of Preparation March 1995

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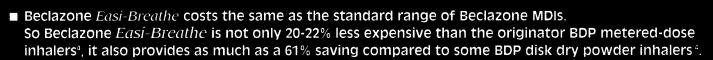
- Lindgren S. Bake B. Larsson S. Eur J Respir Dis 1987;**70**:93-98
 Crompton G.K. Eur J Respir Dis 1982;**63**(Suppl. 119):101-104
- 3 Goodman D.E. et al. Am J Respir Crit Care Med 1994;150:1256-1261
- 4 MIMS, February 1995
- 5 Data on file, Baker Norton



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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:** Bronchia' asthma where use of a pressurised inna!er 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 while reducing or discontinuing oral glucconflictosteriols, the recommended bose in adults including givenly and children 12 years and older is susually 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 bronch. 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 Purmicort does not affect the apility to drive and use machines. Purmicort Respules can be mixed with 0.9% aline and with solutions of terbutaline, salbutamol, sodium cromoglycate or ipratropium promide. **Side** effects: Mid-irritation in the throat, coughing and hoarseness and oral candid as shave 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 θ_2 agonist (e.g. terbutaine: 5-10 minutes before inhalation of Pulmicort Respules. Avoid in pregnancy. By agonist legiteroutainers-5-10 minutes before inhalation of Pulmicort Respules. Avoid in pregnancy. Pharmaceutical precautions: Store below 30°C. Use within 3 months of opening the foir envelope Protect opened ampoure from ignt. Use within 12 hours of opening. Legal category: POM. Basic NHS price: Pulmicort Respules 0.25 mg/m (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 00°7/0309. Pulmicort Respules 0.25 mg/ml PL 00°7/0310. For further information contact the product licence holder: Astra Pharmaceutica's Ltd. Home Park, Kings Langley, Herts WD4 about 10 states and 10 states are supported by the states are supported by the support of the product licence holder: Astra Pharmaceutica's Ltd. Home Park, Kings Langley, Herts WD4 about 10 states are supported by the support of the s



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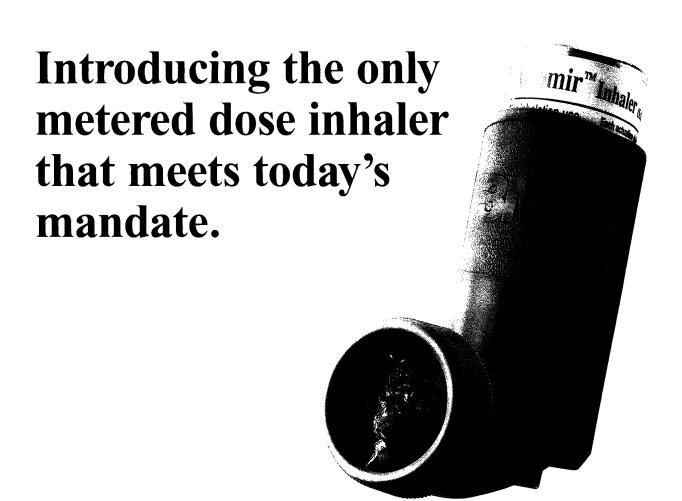


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January 1995

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New Airomir inhaler is the first ever CFC-free metered dose inhaler for asthma—and the only metered dose aerosol inhaler to meet this important initiative.

Airomir inhaler delivers salbutamol sulphate, and has comparable efficacy and safety to the brand leading CFC-salbutamol inhaler^{1,3} at a comparable price.⁴

Switch your asthmatics to Airomir inhaler today, and help make a world of difference. Airomir (salbutamol sulphate inhaler)

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CFC FREE SYSTEM

ARREVIATED PRESCRIBING INFORMATION: Presentation: A pressurised inhalation aerosol delivering Salbutamol Sulphate Ph Eur equivalent to salbutamol 100 mcg into the mouthpiece of the adaptor. Airomir inhaler contains a new propellant, HFA-134a, and does not contain chlorofluorocarbons (CFCs). Indications: For the treatment of reversible airways obstruction associated with asthma, chronic bronchitis or emphysema. It may also be used prophylactically for the treatment of exercise induced asthma. Dosage: Adults and elderly: One or two inhalations as a single dose for the relief of reversible airways obstruction associated with asthma. bronchitis or emphysema. For the prevention of exercise induced asthma, two inhalations prior to exercising. Children: One inhalation for the relief of asthma. increasing to two as a single dose if necessary. One inhalation prior to exercise, increasing to two if necessary. Maximum dose for all patients - eight inhalations in 24 hours. Contraindications: Hypersensitivity to salbutamol or any of the inactive ingredients in the Airomir inhaler. It should not be used in the management of premature labour and threatened abortion. Precautions: Administer cautiously to patients with thyrotoxicosis. Potentially serious hypokalaemia has been reported in patients taking beta-2 agonist therapy. Patients should be advised to seek medical advice if treatment ceases to be effective and/or their asthma seems to be worsening. Patients should not increase the dose without seeking medical advice. Salbutamol and non-selective beta-blockers should not usually be prescribed together. Side-effects: Mild tremor, headache, tachycardia, palpitations, transient muscle cramps. Paradoxical bronchospasm and potentially serious hypokalaemia have been reported in patients taking beta-2 agonists. Pregnancy: There is no experience of Airomir inhaler in human pregnancy. The safe use of salbutamol during pregnancy has not been established but it has been in widespread use for many years without apparent ill consequence. Studies of propellant HFA-134a in pregnant rats or rabbits have not shown any special hazard. Lactation: It is not known whether salbutamol or propellant HFA-134a are distributed into human breast milk. Pharmaceutical precautions: Store below 30°C protected from frost and direct sunlight. As the vial is pressurised no attempt should be made to puncture it or dispose of it by burning. Basic NHS price: £2.30. Product licence number: Pl.0068/0165. Legal Category: POM. Date of preparation: March 1995. References: 1. Data on file. 3M Health Care, Study 1012-SILV, 2. Data on file, 3M Health Care, Study 1037-SILV, 3. Data on file, 3M Health Care. Study 1031-SILV. 4. MIMS March 1995. Date of preparation of literature: March 1995. Further information is available from the 3M Health Care Information Scientist: Telephone Loughborough (01509) 611611. Pharmaceutical Division. 3M Health Care, Loughborough, England, 3M and Airomir are trademarks of the 3M Company.



ion Kit for details.) 1g to d by the i.p. route Ldialysis fluid. Most b.i.d.; severe infections, ty. Cystic fibrosis - up y in three divided doses. ld be reduced when r filtration <50ml/min. An Ig loading dose may be given with d renal insufficiency. Elderly: sage should not normally exceed mates/Infants/Children: (See Data et for details.) Up to two months: 25 o 60mg/kg/day as two divided doses Over two months: 30 to 100mg/kg/day as wo or three divided doses. Cystic throsis, meningitis, immunocompromised: ip to 150mg/kg/day (max 6g daily) in three divided doses. Sensitivity results meningitis monotherapy. The Infusion kit, in the dosage presented, may not be uppropriate for use in children. ire recommended before commencing ntra-indication Known ersensitivity to cephalosporins. tions Previous anaphylactic iction to penicillin. Administer with aution in early pregnancy, infancy and with concurrent nephrotoxic drug ent. Fortum is excreted in man milk in low concentrations light interference with copper eduction methods may occur. ortum and aminoglycosides hould not be mixed in he same giving set or yringe. Prolonged use ay cause overgrowth of non-susceptible rganisms (e.g. andida. interococci)

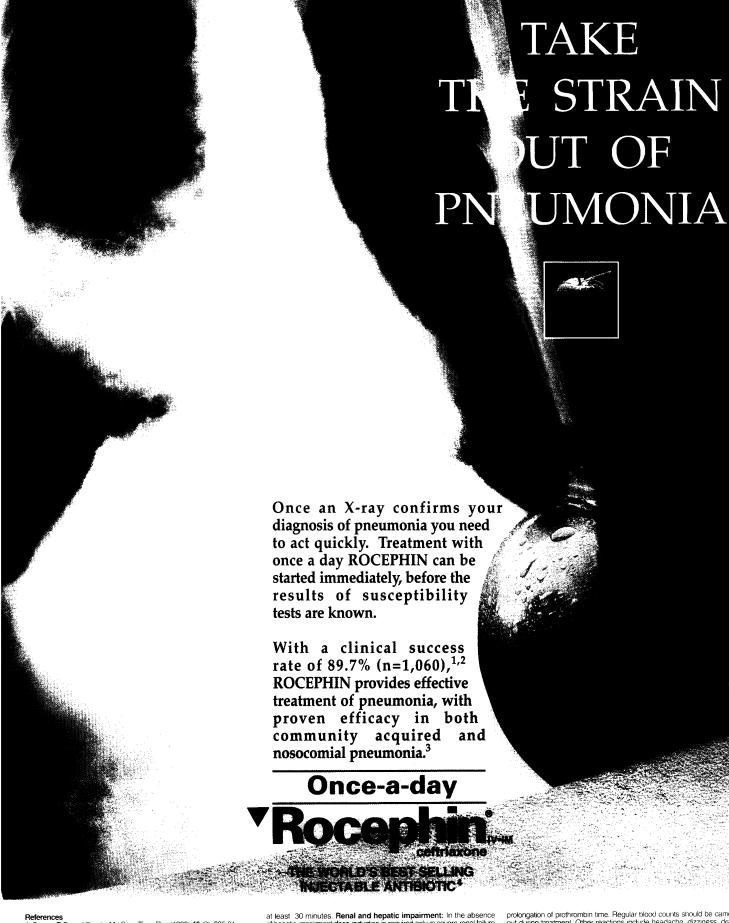
interruption of treatment or other measures. Side effects Adverse reactions occur infrequently; pain and/or inflammation (i.m.) and bitis and/or thrombophlebitis (i.v.), ashes, fever, pruritus, anaphylaxis, GI listurbances, headache, dizziness, araesthesia and bad taste. Transient changes in laboratory values may occur: cosinophilia, positive Coombs' test, hrombocytosis, leucopenia, neutropenia, hrombocytopenia and slight rises in epatic enzymes. Basic NHS cost Packs vials for injection (5 x 250mg, 5 x 00mg, 5 x 1g, 5 x 2g), and an infusion ack (5 x 2g); £9.90 per gram. ortum/Saline Infusion Kit; £20-82. roduct licence numbers Fortum for mection: 250mg: 0004/0304; 500mg: 0004/0292; 1g: 0004/0293; 2g: 0004/0294 Sodium Chloride Intravenous Infusion: 3460/0015. Product licence holder ortum for Injection: Glaco Opera UK Limited, Greenford, Middles UB6 0015. Sodium Chloride Indexenou Infine In: Galen Research Ltd, Craigavon,

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nentremes 1. Brown, R.B. and Sands, M., Curr. Ther. Res (1969), 46 (2), 285-91. 2. Data On File. (GCR B-116 232). 3. Niebuhr, H. *et al.* Chemotherapie Journal (1993), 2, 28-35. 4. Estimaled current cash annual sales worldwide. - Data on File. Roche Products Ltd.

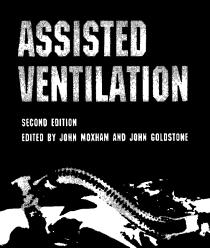
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Brief Prescribing Information
Indications: Pneumonia septicaemia: meningitis: bone, skin and soft tissue
infections: infections in neutropenic patients: gonorrhoea: peri-operative
prophylaxis of infections associated with surgery. Treatment may be started
before the results of susceptibility tests are known. Dosage and
Administration: Rocephin should be administered by deep intramuscular
injection, slow intravenous injection, or as a slow intravenous infusion after
reconstitution of the solution. Adults and children 12 years and over:
Standard dosage -1g once daily. Severe infections - 2-4g normally once
daily. Duration of therapy vanes according to course of disease Gonorrhoea
single dose of 250mg in. Pen-operative prophylaxis - usually single dose
of 1g, colorectal surgery 2g in conjunction with a suitable agent against
anaerobic bacteria. Children under 12 years: Stanoard dosage 20-50mg/kg once daily. Severe infections - maximum 80mg/kg once daily.
Doses of 50mg/kg or over should be given by slow intravenous infusion over

at least 30 minutes. Renal and hepatic impairment: In the absence of hepatic impairment dose reduction is required only in severe renal failure (creatinine clearance <10ml/min), when the daily dose should be 2g or less. No dose reduction is required in liver damage provided renal function is intact. In severe renal impairment accompanied by hepatic insufficiency the plasma concentration should be determined at regular intervals and dosage adjusted. Serum concentrations should be monitored in dialysis. Contra-indications, Warnings etc. Cephalosporin hypersensitivity. Premature infants. Full-term infants during first six weeks of life. Safety in pregnancy has not been established. Precautions: Stated dose should not be exceeded. Caution in patients with a history of hypersensitivity respecially anaphylactic reaction to penicilins or other non-cephalosporin beta-lactam antibiotics. Anaphylactic shock requires immediate countermeasures. Severe renal impairment accompanied by hepatic insufficiency (see Dosage). Side-effects and Adverse Reactions: Gastrointestinal side-effects including loose stools. diarrhoea. nausea vomiting, stomattis and glossits. Cutaneous reactions including maculopapular rash pruritus. urticaria. oedema and erythema multiforme. Haematological reactions including anaemia (all grades). Jeucopenia. neutropenia. thrombocytopenia. eosinophilia. agranulocytosis positive Coombs test and

prolongation of prothrombin time. Regular blood counts should be carried out during treatment. Other reactions include headache, dizziness, drug tever and transient elevations in liver function tests. Rarely, glycosuna oliguria, haematuria, anaphylaxis and bronchospasm. Very rarely, fever and transient elevations in liver function tests. Rarely: glycosuna oliguria, haematurna, anaphylaxis and bronchospasar. Very farely preopitation of ceffinaxone calcium salt in urine in patients on higher than recommended dose. Reversible preopitates of calcium ceffinaxone have been detected by gallibladder sonograms. In symptomatic cases (which are rare). Conservative non-surgical management is recommended Superinfections with yeasts, fungi or other resistant organisms. Rare instances of pseudomembranous colitis. Injection site pain and local phlebitis. Legal Category: POM Presentations and Basic NHS Cost: 250mg vials im. and iv. (containing 250mg ceffinaxone) - £2.87. 1g vials im. and iv. (containing 150mg ceffinaxone) - £11.46. 2g vials for infusion (containing 250mg vials). PL.0031.0171 (1ig vials). PL.0031.0172 (2g vials) Product Licence Numbers: PL.0031.0171 (1ig vials). PL.0031.0172 (2g vials) Product Licence Holder: Roche Products Limited. PO Box 8. Welwyn Garden City. Hertfordshire. AL7.3AY. Full prescribing information is available on request.

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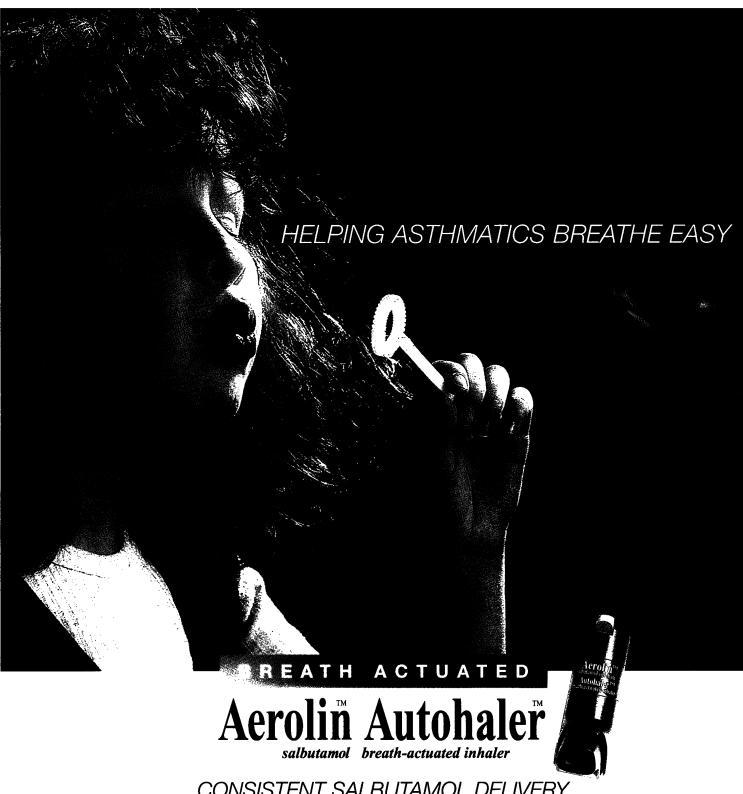
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prescribed together. Side-effects: Mild tremor, headache, tachycardia, palpitations and transient muscle cramps may rarely occur. Paradoxical bronchospasm and potentially serious hypokalaemia have been reported in patients taking B₂ agonist therapy. Pregnancy: The safe use of salbutamol during pregnancy has not been established. Lactation: It is not known whether salbutamol is distributed into breast milk. Pharmaceutical precautions: Store below 30°C protected from frost and direct sunlight. As the vial is pressurised no attempt should be made to puncture it or dispose of it by burning. Basic NHS price: £10.51. Product licence number: PL 68/0117. Legal Category: POM. Date of preparation of advertisement: February 1995. Further information is available from the 3M Health Care Information Scientist: Telephone Loughborough (01509) 611611. Pharmaceutical Division, 3M Health Care, Loughborough, England. Aerolin, Autohaler and 3M, are trade marks of the 3M company.

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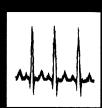
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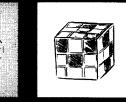
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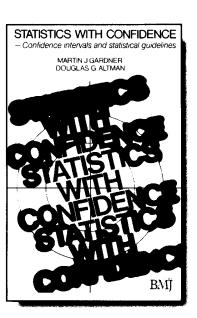
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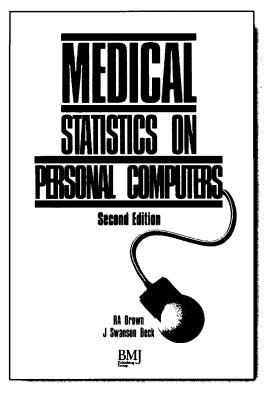
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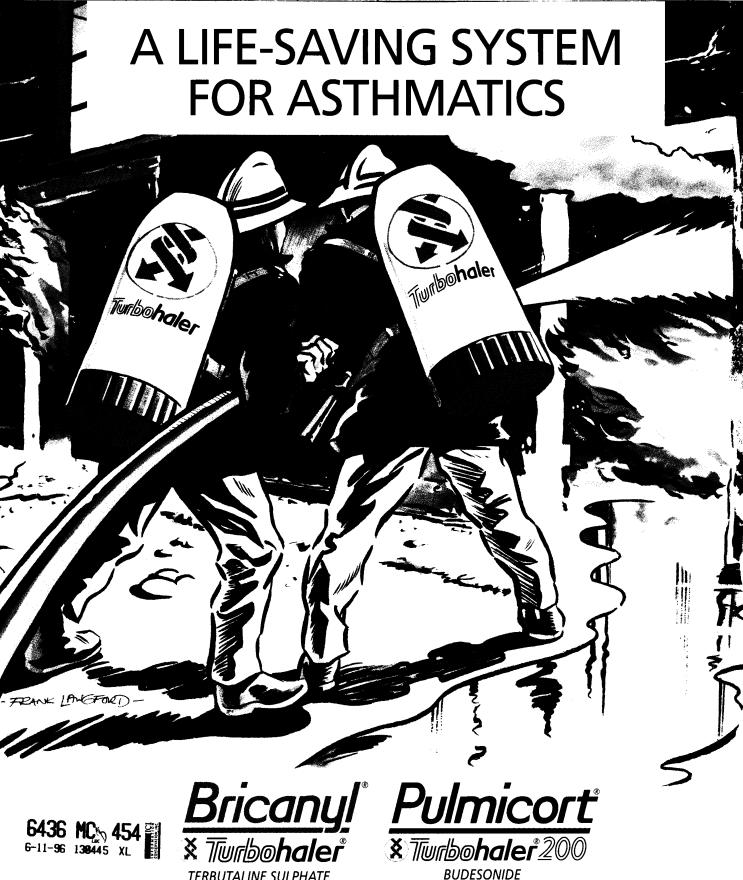
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TERBUTALINE SULPHATE

Abridged prescribing information: Presentation: Bricanyl Turbohaler. Dry powder inhaler delivering 0.5 mg terbutaline sulphate per actuation. Uses Relief and prevention of bronchospasm obstruction is a complicating factor. **Dosage and administration:** Adults and children (including elderly): One inhalation (0.5 mg) as required. Not more than 4 inhalations/day. **Contra-indications, warnings, etc:** Sensitivity to terbutaline sulphate. Precautions: Care should be taken in patients with myocardial insufficiency or thyrotoxicosis. Additional blood glucose measurements are initially recommended in diabetic patients. If treatment becomes less effective or shorter acting, the patient's general condition should be reviewed. Do not use in patients with hypertrophic cardiomyopathy. Potentially serious hypokalaemia may result from B2-agonist therapy. Administer with caution during the first trimester of pregnancy. Do not administer concurrently with nonselective B-blockers. Use with caution with other sympathomimetics. Side effects: Tremor, tonic cramp and palpitations are all characteristic of sympathomimetic amines. A few patients feel tense. **Basic NHS price**: Bricanyl Turbohaler (100 doses) £7.96. **Legal category**: POM. **Product licence** number: PL 0017/0241.

Presentations: Pulmicort Turbohaler 100. 100 μg/puff budesonide dry powder inhaler containing 200 doses. Pulmicort Turbohaler 200. 200 μg/puff budesonide dry powder — Astra Pharmaceuticals — Kings Langley, Herts WD4 8DH. @Registered trade mark

inhaler containing 100 doses. Pulmicort Turbohaler 400. 400 μg /puff budesonide dry power pow inhaler containing 50 doses. **Uses:** Bronchial asthma. **Dosage and administration:** Individual dose. Adults: 200-1600 µg daily in divided doses. Children: 200-800 µg daily in divided do. Maintenance: Use lowest possible dose. Brush the teeth and rinse the mouth out with water all each use. **Contra-indications, warnings, etc:** Active pulmonary tuberculosis. Special care needed in patients with fungal and viral infections in the airways. Avoid administration duri pregnancy. A short course of oral steroids in addition to Pulmicort may be required in patients w excessive mucus in the bronchi. Transfer of patients dependent on oral steroids to treatment w Pulmicort demands special care. See data sheet for further details. Side effects: Mild irritation the throat, hoarseness and oral candidiasis occur in some patients. Rare cases of cataract have been the throat, noarseness and oral candidasis occur in some patients. Rare cases of cataract nave been reported after prolonged use of corticosteroids. Legal category: POM. Basic NHS price: Pulmicor Turbohaler 100 (200 doses) £18.50. Pulmicort Turbohaler 200 (100 doses) £18.50. Pulmicor Turbohaler 400 (50 doses) £18.50. Product licence numbers: Pulmicort Turbohaler 10 (PL 0017/0319 (100 μg/puff). Pulmicort Turbohaler 200 PL 0017/0272 (200 μg/puff). Pulmicort Turbohaler 400 PL 0017/0271 (400 μg/puff). Further information is availated from the product licence holder: Astra Pharmaceuticals Limited, Home Par