bactopoumon

COMPOSITION:
Per 5ml:

Sulfamethoxazole (I.N.N)	400 mg
Trimethoprim (I.N.N)	80 mg
Fluid extract of Tolu	75 mg
Bromhexine (I.N.N) (Hydrochloride)	2 mg

Excipients. Sucrose, 2,0 g; Ethanol, 70,5 mg; Sodium Saccharin, 2,0 mg; Mycrocrystalline cellulose, Strawberry essence, Xanthan gum, Polysorbate 80, Cochineal red, Dimethylpolysiloxane, purified water.

PROPERTIES:
The association of a sulfamide, Sulfamethoxazole, and a diaminopyrimidine, Trimethoprim, gives rise to a drug that is effective against a wide variety of germs causing different diseases. Their union with Fluid extract of Tolu (expectorant) and Bromhexine (mucolytic) enables it to exert an aetiological and symptomatic action in bronchopulmonary processes.

INDICATIONS: Infections of the respiratory tract.
Acute bronchitis an re-intensification of chronic bronchitis, pneumonia, infections of the middle ear and sinusitis.
Treatment and prevention of pneumonia due to Pneumocistis carinii.
Any of the infections cited may be diagnosed and its treatment established by the physician, so this drug should never be used on the patient's own initiative.

POSOLOGY AND METHOD FOR ADMINISTRATION
The doses prescribed by the physician should not be modified or discontinued.
- Adults and children aged over 12 years. 10 ml, twice a day .
- Children aged 6 to 12 years: 5 ml, twice a day
- Children aged 6 months to 6 years. 2.5 ml, twice a day.
- Children aged 2 to 5 months 1.25 ml, twice a day.

It is recommended that during this treatment, you drink sufficient quantities of liquids, if possible.

CONTRAINDICATIONS
- This drug may be contraindicated in people sensitive to any of the substances composing it.
- It should not be used in infants who are premature or aged under 2 months.
- It should not be used with proper supervision in severe haematological disorders.

PRECAUTIONS
- It should be used with care in patients with renal or hepatic insufficiency and in those affected by malnutrition or alcoholism
- It should be used with care in patients with severe allergies or bronchial asthma.
- Special care should be taken with treating elderly patients or bronchial asthma. Avoid excessive exposure to the sun or use of sun lamps.

Pregnancy and lactation:
The possible use of this drug in pregnancy and lactation periods should be evaluated and specified by the physician.
Since it contains ethanol as an excipient, it may be a cause of risk in patients with hepatic disease, alcoholism, epilepsy and in pregnant women and children.

WARNINGS
Every 5 ml of this drug contains 2 g of sucrose, witch should be taken into account by diabetic patients.

INCOMPATIBILITIES
This drug may have an influence on the action of others that are being taken simultaneously, so if you are being treated for diabetes, epilepsy or with drugs affecting blood coagulation, you should inform your doctor of this circumstance.

SIDE EFFECTS
This drug is generally well tolerated by most patients
However, in certain cases, nausea, vomiting, diarrhoea or respiratory difficulties may arise.
More seldom redness of the skin may appear and more severe coetaneous eruptions in elderly patients.
Consult our doctor immediately if any of the foregoing symptoms is manifested.

INTOXICATION AND ITS TREATMENT
The physician should be notified of the accidental taking of this drug or overdosage, stating the amount of product ingested. The commonest symptoms of overdosage are vomiting, nausea, dizziness and confusion.
If overdose or accidental ingestion occurs, consult the Toxicological Information Service. Phone. 91 562 04 20.

DOSAGE FORM
Suspension. Bottle containing 100 ml.

MEDICINES SHOULD BE KEPT OUT OF THE REACH OF CHILDRENS

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