MUXOL 0.3 PER CENT, oral solution

COMPOSITION
Ambroxol hydrochloride ..... 0.300 g
quantity corresponding to ambroxol base ..... 0.273 g
For 100ml of oral solution
Excipient: glycérol, sorbitol 70 per cent crystallizable, methyl parahydroxybenzoate, propyl parahydroxybenzoate, banana flavouring, monohydrated citric acid, purified water.

PHARMACEUTICAL FORM
Oral solution (180 ml bottle).

PHARMACO-THERAPEUTIC CLASS
MUCOLYTICS
(R : Respiratory system)

NAME AND ADRESS OF THE PRODUCER
Laboratoires LEURQUIN MEDIOLANUM
68-88 rue Ampère - Z.I. des Chanoux
93330 NEUILLY SUR MARNE - FRANCE
Medical information : ☎ 33 (0)1 49 44 29 29

WHEN TO USE THIS MEDICINE
Treatment of problems with bronchial secretion in adults, notably during acute bronchial illness or acute episodes of chronic bronchopneumonia.
This medicine is an expectorant. It facilitates the evacuation of the bronchial secretions by coughing.

CAUTION!
WHEN NOT TO USE THIS MEDICINE
This medicine MUST NOT BE USED if there is a history of allergy to this medicine or to one of its ingredients.
IF IN DOUBT, YOU MUST CONSULT YOUR DOCTOR OR YOUR PHARMACIST.

SPECIAL WARNINGS
This medication should not be used by patients with rare hereditary intolerances to fructose.

PRECAUTIONS FOR USE
It is advisable not to take any antitussive medicine or any medicine that dries the bronchial secretions during your period of treatment with this medicine.
IF IN DOUBT, DO NOT HESITATE TO ASK YOUR DOCTOR OR PHARMACIST

INTERACTIONS WITH OTHER DRUGS AND OTHER INTERACTIONS
TO AVOID POSSIBLE INTERACTION BETWEEN SEVERAL DRUGS, YOU MUST SYSTEMATICALLY INFORM YOUR DOCTOR OR PHARMACIST OF ANY OTHER TREATMENT THAT YOU ARE RECEIVING.

PREGNANCY - BREAST FEEDING
It is preferable not to use this medicine during pregnancy.
If you discover that you are pregnant during the treatment, consult your doctor who is the only person capable of judging whether you should continue.
In general, during pregnancy or when breastfeeding, you should always ask for your doctor's or your pharmacist's opinion before taking any medicine.

LIST OF EXCIPIENTS TO KNOW FOR A USE WITHOUT ANY RISK FOR SOME PATIENTS
Methyl parahydroxybenzoate, propyl parahydroxybenzoate, glycerol.

HOW TO USE THIS MEDICINE

DOSAGE
RESERVED FOR ADULTS.
The dosage of this medicine is 1 table-spoonful in the morning and evening.

ADMINISTRATION
Oral.

FREQUENCY AND TIMES WHEN THE MEDICINE MUST BE TAKEN
The medicine must be taken at regular intervals.

DURATION OF THE TREATMENT
The duration of the treatment should not exceed 8 to 10 days without medical supervision.

POSSIBLE SIDE EFFECTS
LIKE ANY ACTIVE PRODUCT, THIS MEDICINE MAY CAUSE CERTAIN SIDE EFFECTS IN SOME PEOPLE :
Possible minor gastro-intestinal problems such as nausea, vomiting, abdominal pains which will diminish when the dosage is reduced.
Also reported:
- cases of cutaneous mucous reactions like erhythema, rashes, pruritus, urticaria,
- very rare cases of anaphylactoid reactions with shock and Quincke's oedema, which developed favourably in the cases reported.

In such cases, the treatment must be stopped.
Cases of headaches and vertigo were also very rarely reported.
Owing to the presence of sorbitol, this medication can cause slight digestive disorders (diarrhoea).
Calories: 2.6kcal per gram of sorbitol.
Can cause allergic reactions to methyl and propylparahydroxybenzoates (possibly delayed).

ANY UNWANTED SIDE EFFECTS NOT INDICATED IN THIS LEAFLET SHOULD BE REPORTED TO YOUR DOCTOR OR PHARMACIST.

CONSERVATION
DO NOT EXCEED THE USE-BY DATE ON THE EXTERNAL PACKAGING

DATE OF REVISION OF THIS LEAFLET
April 2008

Laboratoires LEURQUIN MEDIOLANUM

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