

pharbiol succinimide 3g

powder for making a drinkable solution in single-dose sachet

Qualitative and quantitative composition

SUCCINIMIDE 3 g

Excipients: sucrose, anhydrous colloidal silica, lemon essence. Per single-dose sachet

Pharmaceutical form

Powder for making a drinkable solution, box containing 30 sachets

Pharmaco-therapeutic class

For the treatment of hyperoxaluria (G. Urology)

Name and address of marketing authorization holder

S.E.R.P. - Immeuble le Triton
5 Rue du Gabian - 98000 MONACO

WHEN TO USE THIS DRUG

Supplementary treatment for certain forms of renal lithiasis (kidney stones) in adults.

CAREFUL !

When not to use this drug

This drug **MUST NOT BE USED** in patients with a history of allergy against any of its ingredients
IF YOU ARE IN ANY DOUBT, IT IS ESSENTIAL TO ASK YOUR DOCTOR OR PHARMACIST FOR ADVICE.

Precautions

Your diet should be **low in calcium** (avoid milk and dairy products) and oxalates. You should drink large quantities of water (preferably with low mineral content) during treatment.

IF YOU ARE IN ANY DOUBT, IT IS ESSENTIAL TO ASK YOUR DOCTOR OR PHARMACIST FOR ADVICE.

Drug interactions and other interactions

IN ORDER TO AVOID POSSIBLE DRUG INTERACTIONS, ALWAYS INFORM YOUR DOCTOR OR PHARMACIST IF YOU ARE TAKING OTHER DRUGS.

Pregnancy and breast-feeding

This drug should not be taken by pregnant or breast-feeding women.

If you find out that you are pregnant during the course of treatment, CONSULT YOUR DOCTOR because only he or she can decide how your treatment should continue.

HOW TO USE THIS DRUG

Dosage

FOR ADULT USE ONLY

As a general rule, take one sachet, three times a day.

Administration

Oral

The sachet should be dissolved in a large glassful of water (with or without sugar) and taken before a meal.

ADVERSE REACTIONS

THIS DRUG CAN INDUCE UNDESIRABLE ADVERSE REACTIONS IN SOME CASES:

problems affecting the digestive system or skin eruptions which disappear as soon as the drug is discontinued.

DO NOT HESITATE TO ASK YOUR DOCTOR OR PHARMACIST FOR ADVICE AND YOU MUST INFORM THEN IF YOU HAVE AN ADVERSE REACTION WHICH IS NOT MENTIONED IN THIS PACKAGE INSERT.

DO NOT USE AFTER THE EXPIRY DATE WHICH IS CLEARLY MARKED ON THE PACKAGING

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Last updated of the notice: June 2004