

VIBTIL 250 mg ®

Coated tablet

This is a medicine. Read the leaflet carefully. Ask your doctor or pharmacist for advice. Consult your doctor if symptoms persist.

FOR ADULTS ONLY

COMPOSITION

Dry aqueous extract of linden sapwood250mg
Excipients : Saccharose, light magnesium carbonate, calcium carbonate, hydrogenated castor oil, corn starch, microcrystalline cellulose, gum arabic, talc, alkaline methacrylate copolymer (12,5 % solution), titanium dioxide (E 171), gelatine, carnauba wax, white beeswax, synthetic esters of saturated fatty acids and of alcohol, for one coated tablet.

PHARMACEUTICAL FORM

Box of 40 coated tablets.

PHARMACO-THERAPEUTIC CLASS

CHOLAGOGUE-CHOLERETIC. (A : Digestive system and metabolism).

WHEN TO USE THIS MEDICINE

This medicine is traditionally used to facilitate the body's elimination functions.

PRECAUTIONS FOR USE

USE THIS MEDICINE WITH CAUTION in case of biliary tract obstruction or serious liver disease.
IF IN DOUBT, DO NOT HESITATE TO ASK YOUR DOCTOR OR PHARMACIST FOR ADVICE.

List of excipients that need to be known for safe use in certain patients :

- Hydrogenated castor oil,
- Corn starch (can be harmful in case of celiac disease).

WHEN NOT TO USE THIS MEDICINE

IF IN DOUBT, IT IS INDISPENSABLE TO ASK YOUR DOCTOR OR PHARMACIST FOR ADVICE.

DRUG INTERACTIONS AND OTHER INTERACTIONS

IN ORDER TO AVOID POSSIBLE INTERACTIONS BETWEEN SEVERAL MEDICINES, YOU MUST ALWAYS INFORM YOUR DOCTOR OR PHARMACIST OF ANY OTHER TREATMENT UNDERWAY.

HOW TO USE THIS MEDICINE

FOR ADULTS ONLY.

Swallow 1 to 2 tablets with a little water before your 3 main meals or when in discomfort.
Oral route.

UNDESIRABLE EFFECTS

LIKE ANY ACTIVE SUBSTANCE, IN CERTAIN PERSONS THIS MEDICINE CAN PRODUCE SIDE EFFECTS SOME OF WHICH ARE MORE ADVERSE THAN OTHERS : Risk of diarrhoea at high doses.
TELL YOUR DOCTOR OR YOUR PHARMACIST ABOUT ANY UNWANTED OR ADVERSE EFFECT NOT MENTIONED IN THIS INFORMATION LEAFLET.

STORAGE

Do not exceed the expiry date stated on the external packaging.
Store in a dry place.

DATE OF REVISION OF THE LEAFLET

June 2004.

NAME AND ADDRESS OF RESPONSIBLE FOR PLACING MEDICINAL PRODUCT ON THE MARKET

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