

laluset®

CREAM sodium hyaluronate 0.2%

PROPERTIES

laluset® contains sodium hyaluronate, namely the sodium salt of hyaluronic acid.

COMPOSITION

Sodium hyaluronate 0.2%, Cetearyl glucoside, decylester of oleic acid, cetearyl alcohol, sodium lauryl sulphate, sodium cetearyl sulphate, pure anhydrous glycerine, sorbitol 70%, sorbic acid, methyl parahydroxybenzoate, propyl parahydroxybenzoate, trisodic citrate dihydrate, deionized water.

FORMS AND PRESENTATIONS

100 g pressurized bottle 25 g or 100 g tube

ADMINISTRATION ROUTES AND POSOLOGY

Clean the wound thoroughly before each application, then proceed as follows.

Apply a layer of cream about 2-3 mm thick directly onto the wound, or spread the cream onto a sterile dressing and apply it to the wound.

The dressing can be held in place with a suitable bandage.

WARNINGS AND PRECAUTIONS FOR USE

Tube: Tighten the screw-cap carefully to close the tube after each use.

Pressurized bottle: It is recommended to clean the nozzle every new use by removing the small amount of the remaining cream, by briefly pressing the key dispenser.

Attention: Container under pressure; it may burst if heated.

Protect from sunlight and do not expose to temperatures exceeding 50 $^{\circ}$ C.

Keep away from sources of heat, hot surfaces, sparks, naked flames or other sources of ignition.

Do not pierce or burn the spray can, even after use.

Not smoking.

Do not swallow.

Avoid spraying in the eyes.

Follow the appropriate procedures for cleaning and disinfecting wounds, burns and surrounding areas of skin.

The cream left on the wound from the previous application can be easily cleaned away with water before the next application.

The use of laluset® on venous ulcers does not preclude the use of a compression treatment (socks or bandages).



REV.00 Instructions for use Ialuset® cream tube and BOV - June 2020

Do not use laluset® if the packaging is damaged.

If you have any doubts about using laluset® consult your doctor or pharmacist.

For external use only.

Keep out of the reach of children.

CONTRAINDICATIONS

laluset® must not be used in case of known hypersensitivity to sodium hyaluronate, or to one or more of the other components.

ADVERSE REACTIONS

Local hypersensitivity reactions, such as eczema, have rarely been observed.

IF YOU NOTICE AN ADVERSE REACTION WHICH IS NOT MENTIONED IN THIS LEAFLET, ASK YOUR DOCTOR OR PHARMACIST FOR ADVICE.

STORAGE

Do not use after the expiry date shown on the packaging.

This medical device must be stored at temperature below 25°C.

Year of CE certification: 2012

Last patient information leaflet review: June 2020

Shelf life: 36 months.

The expiry date indicates the maximum validity of the medical device

MANUFACTURER

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DISTRIBUTOR

(Name and address of the Distributor)



Protect from sunlight



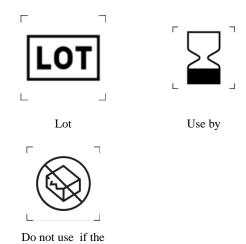
See the instructions for use



Upper limit of temperature



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packaging is damaged