

URGOTUL START



LIPIDO-COLLOID CONTACT LAYER IMPREGNATED WITH NOSF

Indication: Low exuding chronic wounds with risk factors of healing delay

- Inhibits excess proteases
- Promotes the action of growth factors
- Re-balances the environment of chronic wounds
- Painless and atraumatic removal

URGOTUL  START

Supplied in boxes of individually pouched and sterile dressings, ready to use

SIZES

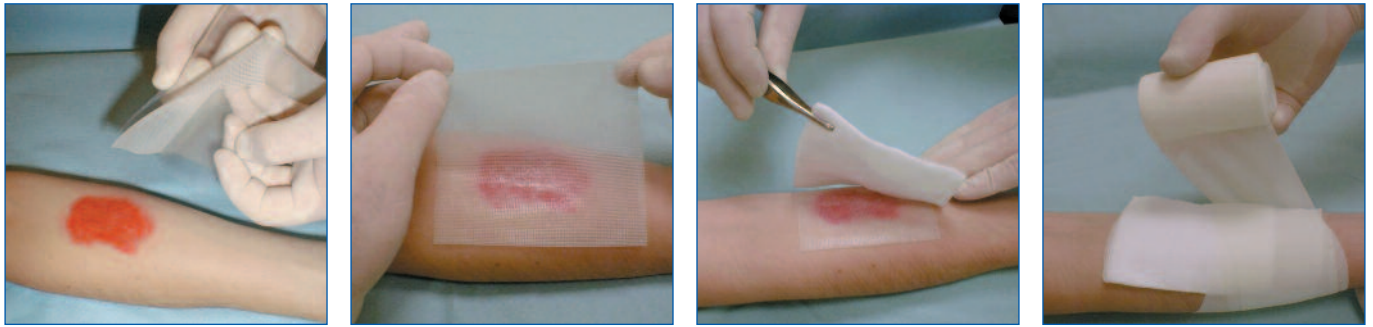
10 x 12 cm

15 x 20 cm

UNITS PER BOX

10 pouches

5 pouches



DESCRIPTION

■ Sterile, non-occlusive, non-adhesive, lipido-colloid contact layer impregnated with NOSF (issued from the Lipido-Colloid Technology, exclusive patented innovation by Urgo Laboratories).

COMPOSITION: **Urgotul® Start** is a lipido-colloid dressing made of a polyester mesh impregnated with hydrocolloid (carboxymethylcellulose), vaseline and NOSF particles (Nano Oligo-Saccharide Factor).

METHOD OF STERILIZATION: Sterilized by ionizing radiation.

PROPERTIES

- In contact with wound exudate, the Lipido-Colloid Technology combined with NOSF provides optimal benefits for the wound healing process:
 - Thanks to the hydrocolloid particles, the Lipido-Colloid Technology forms a gel and creates a moist environment promoting the wound healing process. It allows effective movement of growth factors so that the key cells (fibroblasts, keratinocytes, macrophages) can exert their action in the wound healing process.
 - In contact with wound exudate, NOSF forms a gel which binds preferentially to damaged areas. It interacts with the wound environment by limiting the MMP's activity (Matrix MetalloProteases) which degrade the ECM (extracellular matrix) components. It also promotes the action of the growth factors.
- In contact with wound exudate, the hydrocolloid particles interact with the petroleum jelly component, to form a lipido-colloid gel creating a moist environment favourable for the healing process. The NOSF is maintained in this lipido-colloid gel, providing an activity on contact with the wound only.
- **Urgotul® Start** does not adhere to the wound. Removal of the dressing is painless and atraumatic for the wound.
- Flexible and conformable, **Urgotul® Start** is especially suitable for wounds located in awkward areas.

INDICATIONS

- **Urgotul® Start** is indicated for the treatment of chronic low exuding wounds after debridement, particularly if healing delay is clinically suspected.

DIRECTIONS FOR USE

METHOD OF USE

- Cleanse the wound with saline solution.
- If an antiseptic is first used, rinse the wound thoroughly with saline solution before applying **Urgotul® Start**.
- Remove the protective transparent films from the dressing.
- Apply **Urgotul® Start** directly onto the wound.
- Cover **Urgotul® Start** with a secondary dressing (sterile pad suitable for absorption of exudate).
- Secure **Urgotul® Start** with a retention bandage (Nylex®).
- For the treatment of leg ulcers, a compression system bandage should be applied.
- **Urgotul® Start** should be changed every 3 to 4 days depending on the condition of the wound.
- The minimum period of treatment with **Urgotul® Start** is 4 to 5 weeks.

PRECAUTIONS

- In case of clinical signs of infection, prior treatment of infection is recommended before using **Urgotul® Start**.
- Possible stinging or pain sensations have been reported, especially at the start of the treatment. These are related with the action on the healing process and rarely warrant discontinuation of the treatment.
- **Urgotul® Start** should be stored flat, away from light, heat and moisture.
- Do not use if sterile protection has been damaged.

LEGAL STATUS

 0459 Medical device