

Urgotul[®] Duo



LIPIDO-COLLOID CONTACT LAYER WITH A PROTECTIVE PAD

Indication: Low to moderate exuding wounds

- Healing in a moist environment
- Painless and atraumatic removal
 - Proliferation of fibroblasts
 - Protection of the wound

Urgotul[®] Duo

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

| SIZES | UNITS PER BOX |
|------------|---------------|
| 10 x 12 cm | 10 pouches |
| 15 x 20 cm | 10 pouches |

Urgotul® Duo



DESCRIPTION

- Absorbent sterile non-occlusive, non-adhesive, lipido-colloid dressing (issued from the Lipido-Colloid Technology, exclusive patented innovation by Urgo Laboratories).

COMPOSITION: Urgotul® Duo is a lipido-colloid dressing made of a polyester mesh impregnated with hydrocolloid (carboxymethylcellulose) and vaseline particles, combined with a protective absorbent pad (100% viscose).

METHOD OF STERILIZATION: Sterilized by ionizing radiation.

PROPERTIES

- In contact with wound exudate, the hydrocolloid particles from **Urgotul® Duo** contact layer interact with the petroleum jelly component, to form a lipido-colloid gel creating a moist environment favourable for the healing process.
- Fatty in its chemical composition, without being greasy to the touch, **Urgotul® Duo** does not adhere to the wound or to the surrounding skin. Removal of the dressing is painless and atraumatic for the wound.
- **Urgotul® Duo** promotes fast healing by stimulating fibroblasts proliferation.
- The pad of **Urgotul® Duo** protects the wound and absorbs exudate from the wound. Application is easier, as there is no need to apply a secondary dressing. It is also more comfortable for the patient.
- The transparent films enable to apply **Urgotul® Duo** without touching the mesh and without sticking to latex gloves.
- Flexible and conformable, **Urgotul® Duo** is especially suitable for wounds located in awkward areas.

INDICATIONS

- **Urgotul® Duo** is indicated for the local treatment of acute wounds (burns, skin abrasions, traumatic wounds...) and chronic wounds (leg ulcers, pressure ulcers...) at the granulation/epithelialisation stage.
- **Urgotul® Duo** is also indicated for the treatment of wounds caused by congenital epidermolysis bullosa.

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® Duo**.
- Remove the protective transparent films from the dressing.
- Apply **Urgotul® Duo** directly onto the wound. The TLC mesh should be applied onto the wound side.
- If needed, **Urgotul® Duo** can be cut using sterile scissors to match to the size of the wound.
- Secure **Urgotul® Duo** with a retention bandage (Nylex®) or adhesive plaster.
- **Urgotul® Duo** should be changed every 2 to 4 days, up to 7 days depending on the condition of the wound.
- In case of epidermolysis bullosa, **Urgotul® Duo** should be changed every 1 to 3 days.

PRECAUTIONS

- If clinical signs of local infection appear, the treatment may be continued under medical supervision. However, it is preferable to use **Urgotul® Duo Silver** instead.
- **Urgotul® Duo** should be stored flat, away from light, heat and moisture.
- Do not fold **Urgotul® Duo**.
- Do not use if sterile protection has been damaged.

LEGAL STATUS

CE 0459 Medical device