CREATE A NURTURING ENVIRONMENT FOR SKIN TO RESTORE ITSELF

Acute wounds (burns, traumatic wounds, abrasions, post-operative wounds), chronic wounds (leg ulcers, pressure ulcers and diabetic foot ulcers) and Epidermolysis Bullosa*



SAME GREAT PRODUCT. NEW NAME. For more information visit www.urgomedical.us/transition Urgo Medical NA, formerly SteadMed Medical







WHAT IS UrgoTul?



Jellifying particles

Lipophilic substances

x150 x150 A flexible, non-adherent contact layer featuring the TLC Restoring Matrix that when in contact with wound 20 exudate forms a lipidocolloid gel on the surface of the wound. This gel provides a moist environment 95% optimal for multiplication of key cells patients Report No. RE/DA/2013-164/LAP. for wound healing. With UrgoTul fibroblasts migrate within the wound area, proliferate 70% and synthesize the extracellular greater matrix to form granulation tissue, at 48h⁵ bringing the edges of the wound together.³ Maintaining coating after 24 hours 100 initiale maintaining coating 90 80 0 70 60 50 UrgoTul keeps its integrity over time, 40 without drying out and ensuring atraumatic 30 and pain-free dressing changes.7,8 20 % of i 10 0 UrgoTul **Jelonet**[®] Adaptic Greazy gauze

*95% of patients reported 'no pain' or 'less pain' during dressing changes upon switching to UrgoTul contact layer from another dressing for their acute wound when compared to the period prior to the switch.

UrgoTul FACILITATES THE PROLIFERATION OF KEY CELLS TO RESTORE SKIN



UrgoTul Ensures PAIN-FREE AND ATRAUMATIC DRESSING CHANGES⁴

Character of lesions following dressing removal5



Significant absence of lesions after removal up to 8 days



Eradication of fibroblasts and of the extra-cellular matrix after removal

Clinical consequences at removal



With TLC Restoring matrix (UrgoTul)



With traditional gauze

WHAT DO PATIENTS HAVE TO SAY?



UrgoTul RANGE IS CLINICALLY PROVEN IN MORE THAN 11,600 PATIENTS¹⁰

One-year old child A 13-month-old child presenting with presenting with a partial-thickness a traumatic wound burn on the inside on the forehead of the arm ; after 15 following a road days of treatment traffic accident ; after with UrgoTul, 7 days the wound the wound has fully has healed¹¹ URGO epithelialised¹¹ UrgoTul Can Can Can be used be meshed with other be cut dressing Size with border (inch) Size with border (cm) No. per box Code 10 2×2 5×5 506487 4 x 5 10 x 12 10 506488

Meaume S. Urgotul[®]: a novel non-adherent lipidocolloid dressing. British Journal of Nursing. 2002, Vol 11, N°16. 2. Bernard FX et al. Effects of a lipidocolloid dressing on the production of the extracellular matrix. Journal des Plaies et Cicatrisations, 2007 (study conducted on Urgotul). 3. Bernard FX et al. Stimulation of the proliferation of human dermal fibroblasts in vitro by a lipidocolloid dressing. Journal of Wound Care, May 2005; 14 (5): 215-220 (study conducted on Urgotul). 4. Meaume S et al. The importance of pain reduction through dressing selection in routine wound management: the MAPP study. J Wound Care. 2004;13(10):409-13. 5. Study report No. S/2003-007/BIOalternatives. 6. FX. Bernard et al., Effects of dressings on cellular proliferation and evaluation of their adherence to fibroblasts by in vitro culture. Oral communication, CPC 2005 7. M. Le Berre, Y. Lurton et al., Coated dressings: gauzes/contact layers.
Poster, CPC 2005, Paris. 8. M. Le Berre, Y. Lurton et al., Coated dressings: gauzes/contact layers. Oral communication, CPC 2005, Paris. 9. Benbow M., Iosson, G. A clinical evaluation of UrgoTul to treat acute and chronic wounds. Br J Nurs 2004; 13: 2, 105-109. 10. White, R., Cowan, T., Glover, D. Supporting evidence-based practice: review of TLC healing matrix (2nd edition). MA Healthcare Ltd, London, 2015. 11. A. Le Touze et al., Interest of a new dressing (UrgoTul) in the management of wounds in a pediatric setting: results from a european clinical study. Poster, CPC 2005, Paris.

6 x 8

15 x 20

10

Prior to use, be sure and read the entire Instructions for Use package insert supplied with the product for Device Intended Use, Description, Contraindications, Warnings, Precautions, Adverse Events, and Instructions for Use.



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