### **Technology matters clinically** Absorbs and locks in exudate, even under compression, to reduce risk of leaks and maceration<sup>3,4,9</sup> Efficiently transfers\* exudate away from the wound bed to the secondary dressing<sup>1,2</sup> Promotes autolytic debridement to support clean and easy a clean wound bed upon one-piece removal<sup>3,4,8,9</sup> dressing removal<sup>4</sup>



### With Hydrolock® technology

Unlike traditional gelling fibres, Exufiber® and Exufiber® Ag+ are non-woven polyvinyl alcohol fibre dressings. On contact with exudate they transform into a gel. The tightly packed fibres keep exudate locked in while the capillary action enables transfer of exudate to the secondary dressing<sup>1,2</sup>.

\*For Exufiber® Ag+, when exposed to a flow rate of 0.6ml/h at 40 mmHg pressure for up to seven days.<sup>17</sup>



### **Broad range antimicrobial effect**

Exufiber® Ag+ contains fine silver sulphate crystals. These dissolve on contact with exudate, releasing silver ions, which are proven to kill a broad range of pathogens¹2,¹3,¹4. The antimicrobial effect is rapid (from 3 hours, *in vitro*) and has a sustained effect (up to seven days, *in vitro*)¹2,¹3,¹4. Exufiber® Ag+ prevents biofilm reformation\*\* as part of a biofilm management approach⁵.6.

\*\*As part of a holistic biofilm management approach as per international

### A cost effective approach

Exufiber® dressings help to create an optimal healing environment and reduce the risk of leaks, which means they can be left in place with confidence for up to seven days\*. This promotes undisturbed healing, and can reduce nursing time and costs.

\*Exufiber® and Exufiber® Ag+ can be left in place for up to seven days, depending on wound condition and clinical practice. In addition Exufiber® can be left in place for up to 14 days for donor sites.

|               | Ordering number | Size (cm) | Pcs/RET | Pcs/TRP |
|---------------|-----------------|-----------|---------|---------|
| Exufiber®     | 709900          | 5x5       | 10      | 40      |
|               | 709901          | 10x10     | 10      | 80      |
|               | 709903          | 15x15     | 10      | 60      |
|               | 709905          | 4.5x10    | 10      | 40      |
|               | 709906          | 4.5x20    | 10      | 50      |
|               | 709907          | 4.5x30    | 10      | 60      |
|               | 709904          | 20x30     | 5       | 25      |
|               | 709908          | 1x45      | 5       | 25      |
|               | 709909          | 2x45      | 5       | 25      |
| Exufiber® Ag+ | 603401          | 5x5       | 10      | 40      |
|               | 603402          | 10x10     | 10      | 60      |
|               | 603403          | 15x15     | 10      | 60      |
|               | 603404          | 4.5x10    | 10      | 40      |
|               | 603405          | 4.5x20    | 10      | 50      |
|               | 603406          | 4.5x30    | 10      | 60      |
|               | 603407          | 20x30     | 5       | 20      |
|               | 603400          | 2x45      | 5       | 20      |

Mepilex® Border Flex is the recommended secondary dressing for Exufiber® and Exufiber® Ag+. It combines innovative Flex Technology with our proven Safetac® Technology to create a secondary dressing that stays on and uniquely conforms.

are spent on dressings<sup>15</sup>

References: 1. Mölnlycke Health Care. Data on file. [2018]. 2. Mölnlycke Health Care. Data on file. [2020]. 3. Chadwick P, McCardle J. Open, non-comparative, multi-centre post clinical study of the performance and safety of a gelling fibre wound dressing on diabetic foot ulcers. Journal of Wound Care 2016; 25(4): 290-300. 4. Smet. 5., Beele, H., Saine, L., Suys, E., Henricks, B. Open, non-comparative, multi-centre post market clinical follow-up investigation to evaluate performance and safety on pressure ulcers when using a gelling fibre dressing as intended. Poster Presentation at European Pressure Ulcer Advisory Panel Conference, 2015, Ghent, Belgium. 5. Gille tal. 2017. Evaluation of a Gelling Fiber Dressing with Silver to Eliminate MRSA Biolifilm Infections and Enhance the Healing. Poster presented at the Symposium on Advanced Wound Care Spring meeting/Wound Healing Society (WHS) Annual Meeting 2017, Apr. 05 - 09, 2017, San Diego, CA, USA. 6. Davis, S. C., Li, J., Gil, J., Head, C., Valdes, J., Gilnos, G. D., Solis, M., Higa, A. and Pastar, I. [2019]. Preclinical evaluation of a novel silver gelling fiber dressing on Pseudomonas aeruginosa in a porcine wound infection model. Wound Rep Reg., 27:360-365.
7. Bjarnsholt T, Eberlein T, Malone M, Schultz G. Management of wound biofilm Made Easy. London: Wounds International 2017. 8. Surgical Materials Testing Laboratory. BS EN 13726-1:2002: Test methods for primary wound dressings in Molnycke Health Care. Data on file. [2014.] 9. Davies, P., McCarty, S., An in-use product evaluation of a gelling fibre dressing in wound management. E-poster presentation at Wounds UK Conference, 2017, Harrogate, United Kingdom 10. Molnycke Health Care. Data on file. [2014.] 11. McGrath A [2011] Overcoming the challenge of overgranulation. Wounds UK (71): 42-9-12. Mölnycke Health Care. CE: Performance of Exufiber® Agrin vitro; Antimicrobial effect, silver release kinetics and minimal effective oncentration. [Data on file. 2016.] 13. Hamberg K., Gerner E. and Falkbring S., Mölnycke H

Mölnlycke Health Care AB, P.O. Box 13080, Gamlestadsvägen 3 C, SE-402 52 Göteborg, Sweden. Phone + 46 31 722 30 00 The Mölnlycke, Exufiber, Hydrolock, Mepilex and Safetac trademarks, names and logos are registered globally to one or more of the Mölnlycke Health Care Group of Companies. ©2020 Mölnlycke Health Care AB. All rights reserved HQIM001878. Aquacel and Aquacel Ag Extra are registered trademarks of ConvaTec Inc.

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# Transfers efficiently Removes cleanly





Optimising the space where healing happens



### Chronic wounds present unique clinical challenges

### **Getting the conditions right**

Highly exuding and cavity wounds can be challenging to treat, and painful and worrying for patients. Getting the conditions right for healing is essential. This means a moist environment, without excess exudate. A clean wound bed, undisturbed by slough, residue or debris. And preventing biofilm reformation, which can be a barrier to healing.

### When healing is delayed

Without effective management, wounds can macerate surrounding skin, become infected, or simply refuse to heal. This increases the demands on nurses' time and healthcare providers' costs, and also affects patients' wellbeing, independence and quality of life.

### A fresh take on chronic wounds

The Exufiber® range offers a fresh take on the challenges of highly exuding and cavity wounds. The next generation of gelling fibres aim to optimise the space where healing happens by efficiently\* transferring exudate<sup>1,2</sup> and supporting a clean wound bed³. And Exufiber® Ag+ prevents biofilm\*\*5,6 reformation.

\* For Exufiber® Ag+, when exposed to a flow rate of 0.6ml/h at 40 mmHg pressure for up to seven days. ¹
\*\* Exufiber® Ag+ may be used as part of a biofilm management approach as per international guidelines
[i] a Leapning debridgment & reassessment]

The Exufiber® range addresses the key clinical challenges of highly exuding and cavity wounds

- Exudate pooling
- Slough
- Dressing residue
- Biofilm reformation\*\*

EXUFIBER® AND
EXUFIBER® AG+ HELP
TO CREATE AN OPTIMAL
HEALING ENVIRONMENT

**23**%

MODE OF ACTION

more\*\*\*\* of the exudate absorbed than Aquacel® Extra™8

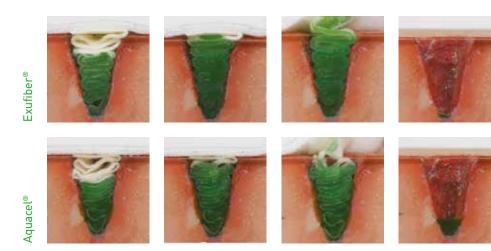
98%

Of clinicians rated Exufiber® as 'easy' or 'very easy' to remove in one piece9

**98**%

Of clinicians reported patient comfort as 'good' or 'very good' for Exufiber®9

### **Proven Transfer Ability**



Watch the full video at Mölnlycke.cor

A cavity model was used to simulate the fluid transfer capability of Exufiber® and Aquacel® ribbon dressings. 5ml of Solution A was added to the cavity and the dressings were left to absorb and transfer. An additional 5ml was then added and the dressings left to absorb and transfer again. Exufiber® demonstrated a better capability to transfer fluid to the secondary dressing than Aquacel®, and when the dressings were removed, less fluid was left in the cavity.

### **Transfers exudate**

Exufiber® dressings transfer exudate efficiently\* from the wound bed to the secondary dressing, locking it in to reduce the risk of pooling, leakage and maceration<sup>3,4</sup>. They can be left in place for up to **seven days**\*\*, allowing undisturbed healing<sup>8,10</sup>.



### Supports a clean wound bed

Residues and debris left in the wound can trigger a foreign body response, and disturb healing<sup>11</sup>. Exufiber® dressings help to break down slough by promoting autolytic debridement<sup>4</sup> They can also be relied upon to stay intact both during use and at removal<sup>3,4,9</sup>.



### Prevents biofilm reformation

Biofilms are present in almost all chronic, non-healing wounds and their presence may prevent healing<sup>7</sup>. Exufiber<sup>®</sup> Ag+ is shown to reduce biofilm bacteria and prevent reformation *in vivo*\*\*\*5,6.

\*For Exufiber® Ag+, when exposed to a flow rate of 0.6ml/h at 40 mmHg pressure for up to seven days.<sup>77</sup>
\*\*Exufiber® and Exufiber® Ag+ can be left in place for up to seven days, depending on wound condition and clinical practice.
In addition Exufiber® can be left in place for up to 14 days for donor sites.

\*\*\*As part of a holistic biofilm management approach as per international guidelines (i.e. cleansing, debridement & reassessment)<sup>7</sup>
\*\*\*\*When comparing lab test results for retention under pressure with Aquacel®, Aquacel® Extra™, Durafiber® and UrgoClean® dressings

### BACKED BY CLINICAL EVIDENCE

### **Outperforms competition**

A recent randomised control trial<sup>16</sup> of 248 venous leg ulcer patients found that Exufiber<sup>®</sup> outperformed Aquacel<sup>®</sup> Extra<sup>™</sup> across multiple measures:

- A positive trend for better wound size reduction
- Clinician satisfaction for overall experience of use, ease of removal, and non-adherance to wound bed
- Clinicians reported better absorption and lock in of exudate, and better lock in of blood and slough

### PATIENT CASE STUDY

Baseline

Exufiber® Ag+

Aquacel® Ag+ Extra

Untreated control

biofilm after treatment

Bacterial counts of

### Manages wound and wound bed

An elderly patient presented with a large, heavily exuding wound on her heel and calcaneous, with approximately 50% sloughy tissue. Initially Exufiber® Ag+ was used as primary dressing to help manage the bioburden and high exudate levels. After two weeks, treatment continued with Exufiber® to manage exudate levels while assisting autolytic debridement. Following 8 weeks of therapy the wound had a 50% area reduction, was moving in positive trajectory and had no clinical signs of infection.

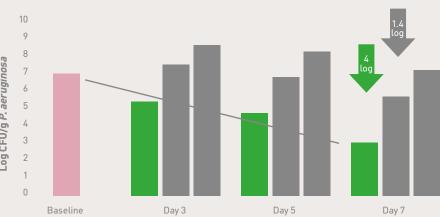




Photographs and case notes kindly supplied by Dr. Paulo Alves, Catholic University of Portugal, Porto, Portugal

## Log CFU/g P. aeruginosa Log CFU/g P. aeruginosa Log CFU/g P. aeruginosa Log CFU/g P. aeruginosa

### Exufiber® Ag+ is superior in reducing biofilm\* bacteria in vivo



Exufiber® Ag+ performs better than Aquacel® Ag+ Extra™ in reducing biofilm bacteria in vivo.

\*As part of a holistic biofilm management approach as per international guidelines (ie cleansing, debridement & reassessment)?

