**PAINLESS AUTOlyTIC DEBRIDEMENT ON 250 CHRONIC WOUNDS BY USING POLYMERIC MEMBRANE DRESSINGS**

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**INTRODUCTION**

Several types of debridement are available for health care professionals but the decision to debride a patients wounds lies on the principle that the healing process needs to be improved and accelerated. Many chronic ulcers are covered with slough and/or necrotic tissue, and are painful and challenging to treat because there is no standard protocol that can be applied to all patients. Very often, these kinds of wounds would need to be debrided more than once. We have noticed that after an initial sharp debridement some wounds became again necrotic/slushy within 24 hours. Optimal and successful wound management needs a treatment plan according to the wound, patients’ parameters and available recourses since a fine line exists between beneficial and harmful debridement. For the past 10 years we have treated these types of ulcers with polymeric membrane dressings (PMDs) due to their unique properties.

PMDs contain a surfactant, superabsorbant starch and glycerol/glyceric, which work synergistically to promote rapid healing and reduction of inflammation. These dressings also facilitate autolytic debridement by loosening bonds between sloughy/necrotic tissue and healthy granulation tissue.

**METHOD AND RESULTS**

The past 10 years we have treated 250 patients with painful necrotic and/or sloughy wounds where autolytic debridement with PMDs was used. A few cases were partially surgically debrided prior to use of PMDs.

PMDs enhance the autolytic process by reducing the interfacial tension level between healthy and necrotic tissue (hence making the necrotic tissue come off easier), and by dissolving dead tissue. Initially the exudates levels increase and daily changes are required to prevent damage to the surrounding skin. Although a slow process, its main advantage over other methods of autolytic debridement is that it’s painless and encourages the healing process by instigating angiogenesis.

Debridement took between 3 to 10 days depending on type and size of the wound.

<table>
<thead>
<tr>
<th>Type of Wound</th>
<th>Total</th>
<th>PMD</th>
<th>Post PMD</th>
</tr>
</thead>
<tbody>
<tr>
<td>Arterial ulcers</td>
<td>26</td>
<td>26</td>
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</tr>
<tr>
<td>Venous ulcers</td>
<td>16</td>
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<tr>
<td>Diabetic ulcers</td>
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<tr>
<td>Mixed ulcers</td>
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<td>42</td>
<td>42</td>
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<tr>
<td>Pressure ulcers</td>
<td>34</td>
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**DISCUSSION**

At our Clinic we focus on continuous debridement, exudate control, prevention of wound trauma and infection as well as pain control. We also take into account changes in the skin such as appearance, structure, mechanical properties and barrier function due to ageing, PMDs help us achieve our goals; as a single treatment modality it’s effective for all phases of wound healing and does not require additional treatments or additives. This made it especially helpful in settings where patients or their family had to perform dressing changes at home.

One reason for primarily choosing autolytic debridement was that surgical intervention was often too stressful for patients with low albumin, HB and iron levels.

As it is impossible to show all cases here I am presenting a representative selection of cases to show how the PMDs work and support our initial decision of using it as a single modality treatment.

**Method:**

On assessment, the wound is a painful, 10 out of 10, covered with yellow slough which could not be removed by surgical debridement due to an arterial occlusion. PMDs were used in order to alleviate pain and instigate autolytic debridement. Analgesia was given every 4 hours.

Pain level reduced from 10 to 6 probably due to PMDs capability to absorb sodium ions by capillary action, from the skin and the subcutaneous tissues resulting in reduced nociceptor conduction, hence alleviating pain. The surfactant dissolves debris by loosening the bonds between the necrotic tissues and absorbing the slough into the dressing. Analgesia was given only at night.

Reinforced autolytic debridement takes place constantly throughout the healing process, clearing the wound from necrotic tissue during dressing wear time. Pain level dropped to zero and no analgesia was needed any more. The dressing does not adhere to the new tissue and therefore does not cause any secondary damage to the wound at dressing changes.

The surfactant reduces surface tension and glycerol helps maintain the moisture level of the wound, together promoting and enhancing a continuous cleansing and debridging of the wound in a painless fashion.

In spite of his poor prognosis and reduced peripheral circulation this wound started to heal.

**Results:**

An 88 year old male with Chronic Obstructive Airway Disease and a huge sacral pressure ulcer caused by spending most of his days sitting upright (to prevent dyspnea). A wound culture showed that the wound was infected with pseudomonas. Silver PMDs moistened with 3 ml normal saline solution were used to hydrate the necrotic tissue, and reduced the infection. No oral antibiotics were prescribed. Pain score 9 at initial assessment.

The surfactant (wound cleanser) and glycerol embedded in the dressing continually help cleanse and debride the wound. The hygroscopic action of glycerin cleans the wound by removing discharge with its osmotic action; reducing edema by inhibiting the lymphatic extension of infection. Pain level dropped to a 4 after two days.

A 92 year old male, diabetic, with a pressure ulcer infected with MRSA. Partial surgical debridement was done in the hospital prior to admittance to our center. Due to extensive bleeding the surgical debridement had to be stopped. She is on antibiotics, Levofloxaxon and Amoxicillin.

**Method:**

PMD is used as a debriding agent. The wound is covered with a thick version of PMD. Over the necrotic tissue we moistened the PMD to enhance the debridement process.

**Results:**

During dressing change a PDM is gently rubbed over the necrotic tissue in a circular motion to enhance the autolytic debridement with a gentle mechanical debridement. Pressure relieving mattress and reposisioning every 2 hours.

Continuous autolytic debridement and healing is evident. The glycerin embedded in PMDs maintains an ideal moisture balance for wound healing; the dressing absorbs exudates and prevents maceration of the periwound area, reduces the interfacial tension between the viable and non-viable tissue, does not adhere to the wound bed, and protects it from further trauma.

A 94 year old man with Alzheimer’s disease with an extensive pressure ulcer. Initial partial surgical debridement was performed in the hospital prior to admittance to our center. Due to extensive bleeding the surgical debridement had to be stopped.

**Method:**

Infected by MRSA after surgical debridement in the hospital. The patients’ pain score was 8 out of 10. The dry sloughy wound was covered by a Silver PMD moistened with 3 ml saline.

**Results:**

Evidence of autolytic debridement. Pain score reduced to 3. The patients’ albumin level was very low and we believe that the availability of glycerin within the PMDs allows it to act as a nutrient and energy substrate, creating a more optimal healing environment that is otherwise almost impossible to achieve in Diabetic patients.

The patient reports no pain at all. The wound is healing nicely and the exudate level is increasing. We no longer need to pre-moisten the dressing prior to application.

A 92 year male, diabetic, with a pressure ulcer infected with MRSA. Previous treatment, i.e. antibiotics and hyaluronic acid cream had no effect on this deteriorating wound.

**Method:**

Infected by MRSA after surgical debridement in the hospital. The patients’ pain score was 8 out of 10. The dry sloughy wound was covered by a Silver PMD moistened with 3 ml saline.

**Results:**

Constant reinforced autolytic debridement of the wound leaves the wound bed, soft, clean, moisturized and smooth, with the surfactant reducing the interfacial tension between the wound and necrotic tissue. Pain score dropped to a level of 2 after a week’s use of PMDs.

**References**

2. Walter JM, Mabbach HJ (2006). Age and Skin structure and function, a quantitative approach (6) protein, glycosaminoglycans, water and lipid content and structure. Skin Research Technology, 12(3) pp 145-154

*PolyMem®* Wound Dressings with and without Silver. Manufactured by Ferris Mfg, 5133 Northeast Parkway, Fort Worth, TX 76106, USA. This case series was unsponsored.