Wound Bed Preparation: Standards of Care and Predictability of the Healing Time of Chronic Wounds

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Abstract

We intend to highlight the need to establish a hypothetical time of healing for chronic wounds. Such innovative aspect within the TIME acronym [tissue, infection/inflammation, moisture balance and edge of wound] is based on a more correct analysis of the general and local symptoms that the patient presents, classifying them in different ways according to their characteristics. The analysis of medical history, pathogenesis and clinical leads to a total score that can direct the clinician to choose principals important and aggressive that can be used according to basic concepts.

Background

To predict the healing time for each patient is not always a simple task. The healing process is the result of a complex interaction between factors related to injury, the treatment used, skills and knowledge of the nursing staff. The best way to shorten the healing time is an accurate staging of the lesion [therefore a proper understanding of the gravity of the wound] and then the decision of the most appropriate therapy. In the last twenty years there have been great strides in the understanding of the biological mechanisms involved in repairing wounds, so now more than ever we have the means necessary to assume a time of complete healing as accurate as possible [1].

Since 2007, Ligresti et al. [2], with the introduction of the TIME - H, relate the healing time with the type of therapy used: it focuses on how the therapy is used with the same type of lesion to make a difference in healing time. Starting in 2008 with the EWMA position document [3] a holistic approach to wound healing is considered but also the complexity of the wounds, the healing process, psychosocial factors involved in healing and economic burden of chronic wounds are considered. Our paper work tends to make available to the medical staff a valid model showing the most correct treatment for a certain type of injury that can allow us to predict the healing time.

Materials and Methods

We performed a study on a group of 40 patients, enlisting 28 females, 12 males aged between 30-87 years. The injuries were wounds and skin ulcers of similar size [50-100cm²], pressure ulcers, venous leg ulcers, diabetic foot, PDS posttraumatic. All patients were treated with one of the principals randomly choosing among those listed with intervals recommended by the type of wound. The study duration was 6 months.

In our proposal we considered a whole TIME and volume to get a severity index [IG], which allows us to have a prediction of healing time. The severity rate also allows us to adopt different therapeutic strategies based on the severity of various injuries. Starting from TIME we built a scale of values that we added from a maximum score of 16 [2-6]. Then, we considered the volume [volume “W” single ulcer or the sum of multiple ulcers]. As shown by Kramer et al. [7] the size and depth of pressure ulcers are good predictors of healing: as lower as the ulcer grade is greater will be the chance of recovery.

We then moved on to the study of pain considering parameter with score 1 as no pain, mild pain score with value 2 [stimulated during the dressing value] and 3 to intense pain [without stimulation] [8]. Finally, we look at general conditions and environmental factors [9-24]. At this point in our algorithm we add the Time and the FAC [factors, anamnestic, social and clinical] and multiply them for pain and volume: the value obtained is our gravity index [IG].

\[
\text{TIME} \times \text{FAC} \times \text{PAIN} \times \text{VOLUME} = \text{IG}
\]

The healing time [TH], as already stated, also depends on the therapeutic strategy undertaken: the numerical value is in direct
relation with the therapy. The TH is=to IG and will not change if the therapeutic choice [soft] resides on the left side of the table; is <30% of IG if therapy undertaken [medium aggressive] is in the middle of the table; is >60% of IG if therapy choice [aggressive] is on the right side of the table. We assume that we are faced with an ulcer from healing [healing time >60 days]. We can change the situation by customizing the type of therapy on the individual case. In this way we can modify the healing time [4].

Take for example a lesion volume between 50 and 100cm3. We can have a lesion T0, I0, M0, E0, thus devoid of necrotic tissue, bacterial contamination, exudate and with re-epithelialisation rate >75%, or a lesion T4, I4, M4 and E4 with 100% of necrotic tissue infections, exudate and the absence of spontaneous re-epithelialization: with the same situation, in any case, the choice of therapy will change the history of the lesion and the timing of healing. For example, the T, we can see how, by choosing a soft treatment [such as autolysis, we have a significant improvement of the lesion as low as 25 days [25-27]. This time can decrease, even drastically, by level up in the type of strategy used: the time passes to 20 days with osmosis, 15 days with the larvae, until a day using hydrotherapy, ultrasound or surgery. Regarding the I: we passed from an improvement of 55 days with saline [strategy blander], to 21 days of antiseptics/ dressings with silver/NPWT [negative pressure wound therapy] up to 7 days after the surgery/ antiseptics/antibiotics/NPWT. The M represents an increase in 55 days with hydrogel/hydrocolloid, hydrofibre 28 days with up to 10 days of the surgery. Finally, the E: re-epithelialization from 100 days with advanced medications, VAC 45 days with up to 10 days with autologus graft [28-56].

**Time of Healing**

The numerical value is in direct relation to the selected therapy. It is=to that of IG patient and its value will not change if the therapy choice [soft] is in the left side of the table therapeutic

It is <30% of IG patient if treatment choice [medium-aggressive] is in the centre of the table therapeutic

It is <60% of IG patient if treatment choice [aggressive] is on the right side of the table.

**Result**

The test results have shown that the margin of error in predicting the healing time was <10% of the 40 patients analysed in detail (Table 1):

**Table 1: Soft: not aggressive treatment.**

<table>
<thead>
<tr>
<th>Type of Wound</th>
<th>P</th>
<th>Performed Treatment</th>
<th>Performed Treatment</th>
<th>Time of Healing</th>
<th>Time of Healing</th>
<th>Difference of Healing in Days Time</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>First 30 days</td>
<td>30-180 days</td>
<td>Expected (days)</td>
<td>Realized (days)</td>
<td></td>
</tr>
<tr>
<td>Post-traumatic wound</td>
<td>40</td>
<td>3=soft</td>
<td>3=MA</td>
<td>21</td>
<td>24</td>
<td>3</td>
</tr>
<tr>
<td></td>
<td></td>
<td>2=MA</td>
<td>2=MA</td>
<td>28</td>
<td>32</td>
<td>4</td>
</tr>
<tr>
<td></td>
<td></td>
<td>2=A</td>
<td>2=A</td>
<td>40</td>
<td>45</td>
<td>5</td>
</tr>
<tr>
<td>Cutaneous vascular</td>
<td>21</td>
<td>12=soft</td>
<td>7=soft</td>
<td>32</td>
<td>35</td>
<td>3</td>
</tr>
<tr>
<td>venous ulcer</td>
<td></td>
<td>7=MA</td>
<td>12=MA</td>
<td>28</td>
<td>30</td>
<td>2</td>
</tr>
<tr>
<td></td>
<td></td>
<td>2=A</td>
<td>2=A</td>
<td>28</td>
<td>32</td>
<td>4</td>
</tr>
<tr>
<td>Diabetic foot</td>
<td>9</td>
<td>4=soft</td>
<td>2=soft</td>
<td>21</td>
<td>24</td>
<td>3</td>
</tr>
<tr>
<td></td>
<td></td>
<td>3=MA</td>
<td>5=MA</td>
<td>30</td>
<td>32</td>
<td>2</td>
</tr>
<tr>
<td></td>
<td></td>
<td>2=A</td>
<td>2=A</td>
<td>40</td>
<td>44</td>
<td>4</td>
</tr>
<tr>
<td>Pressure ulcers</td>
<td>3</td>
<td>1=MA</td>
<td>1=MA</td>
<td>56</td>
<td>59</td>
<td>3</td>
</tr>
<tr>
<td></td>
<td></td>
<td>2=A</td>
<td>2=A</td>
<td>60</td>
<td>65</td>
<td>5</td>
</tr>
</tbody>
</table>

MA: Medium-Aggressive Treatment; A: Aggressive Treatment

Report of the results obtained with the averages of the healing times on a number of 22 patients, obtaining a slightly higher percentage of error of 10% in the prediction of healing time.

**Our protocol**

GI is <23=and ASA 1-5 soft therapy

GI is 24-50 and ASA <4 with infection between 1-2=medium-aggressive therapy

GI is >50 and ASA <4 with infection between 3-4=aggressive therapy

GI is >50 and ASA is 4-5 with infection between 1-2=soft therapy

GI is >50 and ASA is 4-5 with infection between 3-4=medium-aggressive therapy

**Discussion**

Nowadays, various systems today try to give a prediction of healing time, taking into account various parameters. Troxler et al [4] studied the importance of periodic evaluations of the wound, accompanied by measurements of its surface, for the identification of potentially hard-to-heal wounds. The early detection of a reduction
in the size of the wound is set by measuring the progress of the margin [epithelial advancement]. Phillips et al. [57] considering the percentage reduction in venous ulcer area found that in about 77% of cases, healing outcomes could be predicted based on a wound size reduction of more than 44% at three weeks. Margolis et al. [58,59] were able to show that for venous leg ulcers a simple rating system score based on size and duration can give a good indication of the likely outcome at 24 weeks. Falanga et al. [60] incorporated measurement of epithelial advancement into a scoring system on the healing of venous leg ulcers. This system [wound bed score] also examines other characteristics including the extent of skin dermatis around the wound, the presence of eschar; callus and/or fibrosis around a wound, pink or red wound bed, exudate and the volume of the edema (Table 2).

**Table 2.**

<table>
<thead>
<tr>
<th>Protocol A1 30 Days</th>
<th>Soft</th>
</tr>
</thead>
<tbody>
<tr>
<td>T=fibrin</td>
<td>autolytic cream</td>
</tr>
<tr>
<td>I=absent</td>
<td>saline or ringer Lactate</td>
</tr>
<tr>
<td>M=absent</td>
<td>hydrogels or hydrocolloids</td>
</tr>
<tr>
<td>E=depth 0</td>
<td>hydrocolloids, hydrofibres, collagen, ac. Ialuronico</td>
</tr>
</tbody>
</table>

The complexity of the wound is likely to exert a significant influence on the progression of the healing process, and the factors that combine to determine it can be classified into four main groups: patient factors, factors related to the wound, knowledge of the HCP, factors and resources related to the treatment. In a study by Margolis et al. on a group of patients with venous ulcers, it has emerged a correlation between some specific characteristics of the wound and the healing process: wound duration, size, depth of the wound]. Ulcer size [>2cm2], the duration [>two months] and depth

**Table 3.**

<table>
<thead>
<tr>
<th>Protocol A2 60 Days</th>
<th>Moderate</th>
</tr>
</thead>
<tbody>
<tr>
<td>T=slough</td>
<td>autolytic or osmotic dressing</td>
</tr>
<tr>
<td>I=contamination</td>
<td>antiseptic</td>
</tr>
<tr>
<td>M=absent</td>
<td>collagen or hyaluronic acid</td>
</tr>
<tr>
<td>E=depth 0,5</td>
<td>growth factors</td>
</tr>
</tbody>
</table>

For the physiological nature of the healing process, it is inevitable that large wounds will require more time to heal than smaller wounds. In addition, the longer a wound remains open, the greater the risk of complications, such as infections, is present. Therefore, a treatment that reduces the size of the wound and the infection risk is able to offer potential benefits. The presence of necrotic tissue in a wound has been for a long time considered an obstacle to the evaluation of the lesion, as well as a potential predictive factor of delayed healing and a possible outbreak of infection (Table 4).

**Table 4.**

<table>
<thead>
<tr>
<th>Protocol B 120 Days</th>
<th>Aggressive</th>
</tr>
</thead>
<tbody>
<tr>
<td>T=infected surface, static edges</td>
<td>Surgical debridement</td>
</tr>
<tr>
<td>I=contamination</td>
<td>antiseptic, Ag dressing, NPWT, antibiotics</td>
</tr>
<tr>
<td>M=+++infected</td>
<td>polyurethane-NPWT therapy</td>
</tr>
<tr>
<td>E=depth 1-2</td>
<td>allografts, bioengineered implants, dermal Substitutes</td>
</tr>
</tbody>
</table>

In chronic wounds, there is a tendency for the inflammatory response [which is an important element of the initial response to the lesion]. This results in increased production of pro-inflammatory cytokines, reactive oxygen species and proteolytic enzymes [such as certain MMPs, elastase and plasmin]. This activity is combined with a minor issue, for example, inhibitors TIMP (Table 5), and is further enhanced by alterations of pH at the level of the wound bed. Excessive activity of these enzymes causes not only deleterious extracellular matrix destruction, but also inactivation of growth factors. There is a correlation between the state of chronic inflammation of a wound, the high levels of protease exudate and the slowdown in the process of tissue repair. The control of inflammation and the concentration of MMPs [metalloproteases][penetration through exposed tendon, ligament, bone or joint] are essential, as the protease, not only degrade the fabric, but lead to malnutrition of the growth factors [61-69].

**Table 5.**

<table>
<thead>
<tr>
<th>Protocol C1 180 Days</th>
<th>Aggressive with Hospitalization</th>
</tr>
</thead>
<tbody>
<tr>
<td>T=deep necrosis</td>
<td>Surgical debridement</td>
</tr>
<tr>
<td>I= certain infections</td>
<td>H2O2 (abscesses), antiseptics, surgical debridment, dressing Ag, NPWT, antibiotics</td>
</tr>
<tr>
<td>M=+++ infection</td>
<td>surgery, NPWT, polyurethane</td>
</tr>
<tr>
<td>E=depth 2</td>
<td>stimulators or dermal substitutes dermal surgery</td>
</tr>
</tbody>
</table>

In chronic wounds, there is a tendency for the inflammatory response [which is an important element of the initial response to the lesion]. This results in increased production of pro-inflammatory cytokines, reactive oxygen species and proteolytic enzymes [such as certain MMPs, elastase and plasmin]. This activity is combined with a minor issue, for example, inhibitors TIMP (Table 5), and is further enhanced by alterations of pH at the level of the wound bed. Excessive activity of these enzymes causes not only deleterious extracellular matrix destruction, but also inactivation of growth factors. There is a correlation between the state of chronic inflammation of a wound, the high levels of protease exudate and the slowdown in the process of tissue repair. The control of inflammation and the concentration of MMPs [metalloproteases] are essential, as the protease, not only degrade the fabric, but lead to malnutrition of the growth factors [61-69].

**Table 6.**

<table>
<thead>
<tr>
<th>Protocol C2 &gt;180Days</th>
<th>Containment</th>
</tr>
</thead>
<tbody>
<tr>
<td>T= necrosis</td>
<td>enzymatic dressing</td>
</tr>
<tr>
<td>I= certain infections</td>
<td>H2O2 (racolite), Antiseptic, Medicazioni Ag, Antibiotic</td>
</tr>
<tr>
<td>M=+++ infected</td>
<td>Hydrofibres + Ag, Polyurethane</td>
</tr>
<tr>
<td>E=depth 0-2</td>
<td>Hydrocolloids, hydrofibres, hyaluronic acid, growth factors</td>
</tr>
</tbody>
</table>
Gjødsbøl et al. [70] found a significant link between diversity and the density of the bacterial species detected on the diagnostic buffer and the time required for wound healing. Also, the presence in a wound of specific bacterial species has been put in relation with the outcome of healing. For example, the presence of *Pseudomonas aeruginosa* in venous leg ulcers can delay healing. According Mogford et al. [71], an ischemic wound is probably the most common cause of non-healing. Because of a poor perfusion, metabolic gas exchange at the level of tissues become inefficient. It has been shown that the healing of a wound following surgery is compromised by dehydration and by a low body temperature of the patient, factors that are associated with reduced perfusion tissue and poor oxygenation (Table 6).

Physical factors such as diabetes mellitus, obesity, malnutrition, advanced age [over 60], decreased perfusion, peripheral vascular disease, cancers, organ failure, sepsis, and even restrictions mobility, can affect the healing process. Marston et al. [72] have found that improved glycemic control has a positive influence on the outcome in diabetic foot wounds, particularly when dermal substitutes are used. Terms of immunodeficiency, use of immunosuppressive drugs [corticosteroids, azathioprine or methotrexate] [73-75] or the presence of diseases [such as diabetes mellitus] [76-79] known to affect the imuno-inflammatory response, are all circumstances that may influence negatively healing and increase the risk of wound sepsis.

It was also found that psychosocial factors [80-82], such as social isolation, gender, smoking, the economic conditions and the experience of pain could influence wound healing. Stress and depression have been linked to changes in immune function and may therefore adversely influence a wide range of physiological processes, including wound healing. In a human experimental model, it was found that stress and depression had a possible role in the modulation of matrix metalloproteinases [MMPs] and expression of tissue inhibitors of metalloproteinases [TIMP]. According to some studies also the ability to cope with stress is a role in the modulation of matrix metalloproteinases [MMPs] and expression of tissue inhibitors of metalloproteinases [TIMP].

It describes how the therapy used should change significantly and according to the same applies a type of therapy.

Although the feeling of helplessness experienced by some patients, many of them make every effort to ensure that the care they receive meets their needs. Some patients become experts in their own condition, often using the Internet to gather information on it. When a wound is located on a pressure-bearing surface or a mobile area such as around a joint, the choice of the material of the dressing and the method of attachment is of extreme importance. However, Chipchase et al. [84] observed that, while the overall healing rates of foot ulcers were similar, lesions located in the heel tended to heal more slowly. The authors concluded that the outcome was generally favorable, with 65.6% of heel ulcers healed in a median time of 200 days.

Harding [85] has the potential effect of fibroblast senescence on chronic wound healing. There was a correlation between the ratio of senescent fibroblasts/non-senescent fibroblasts and healing outcomes: an accumulation of more than 15% senescent fibroblasts is considered the threshold beyond which wounds will have trouble healing. Moreover, the response to treatment can be an indicator of tissue viability and healing potential. For example, it was suggested that a reduction in wound area of around 15% within one to two weeks of topical negative pressure therapy is a positive indicator of the likely evolution of the wound, and that this observation can be the decision to continued therapy.

As seen above, many attempts and proposals tend to quantify the time required for wound healing. On the basis of previous attempts, our aim is to propose an algorithm not only to accurate the quantification of healing time, but also a precise protocol of treatment to be associated with any situation and any type of wound. It is therefore clear; as the challenge of healing of a skin lesion, cannot be achieved by a single specialist. It's fundamental the collaboration between various specialists. It's obvious that the planning therapy must be organised by a team leader to coordinate the rest of the medical staff [various specialists including plastic surgeon, vascular surgeon, the internist, the nutritionist, the endocrinologist, the dermatologist, etc.] and not by hospital nurses or domiciles medical figures. Only with careful collaboration and clearly careful training you can pursue the goal in the shortest time possible and with a greatest patient comfort.

This protocol sets itself as a general guideline of the ulcer treatment for all types of professionals involved in the management of the patient but is particularly useful for those who approach the world of vulnology [wound treatment], but who do not yet have a detailed knowledge about. It appears a useful tool, since for the first time, it offers a mathematical model in which a score is calculated and according to the same applies a type of therapy.

**Conclusion**

The algorithm that we propose is a useful tool for staging the severity of injuries and provides a simple means to adjust therapy. It describes how the therapy used should change significantly according to the type and severity of the wound that we are facing and how this in turn can affect significantly the healing time estimated. It’s clear that, not always, the mathematical calculations are an exact prediction, but allow however to have a prediction of the initial situation. There are margins of error, especially when it takes over factors that complicate the wound management. It is highly appreciated that, to predict the outcome of recovery in individual patients, it will be necessary to use not a single marker, but the data resulting from a combination of several markers, as we proposed. Many authors have stressed the importance of training the nursing staff, designed to provide the knowledge and skills necessary to establish appropriate treatment and process protocols and forms relating to wound care [56].
However, we can conclude by saying that we do not presume to have created a perfect model for the treatment of skin lesions, we only want to provide a useful tool to the caregiver based on current knowledge and on current therapeutic strategies. We are still waiting for new knowledge that can lead us to an even higher level in the treatment of this complex disease.

References

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