Management of Chronic Non-healing Wounds by Hirudotherapy

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ABSTRACT

A chronic wound is a wound that does not heal in an orderly set of stages and in a predictable amount of time or wounds that do not heal within three months are often considered chronic. Chronic wounds often remain in the inflammatory stage for too long and may never heal or may take years. Chronic wound patients often report pain as dominant in their lives. Persistent pain is the main problem for patients with chronic ulcers. Many wounds pose no challenge to the body’s innate ability to heal; some wounds, however, may not heal easily either because of the severity of the wounds themselves or because of the poor state of health of the individual. Any wound that does not heal within a few weeks should be examined by a healthcare professional because it might be infected, might reflect an underlying disease.

KEYWORDS

Hirudotherapy; Wound; Chronic; Healing

INTRODUCTION

A chronic wound is a wound that does not heal in an orderly set of stages and in a predictable amount of time or wounds that do not heal within three months are often considered chronic. Chronic wounds often remain in the inflammatory stage for too long and may never heal or may take years. Chronic wound patients often report pain as dominant in their lives. Persistent pain is the main problem for patients with chronic ulcers. Many wounds pose no challenge to the body’s innate ability to heal; some wounds, however, may not heal easily either because of the severity of the wounds themselves or because of the poor state of health of the individual. Any wound that does not heal within a few weeks should be examined by a healthcare professional because it might be infected, might reflect an underlying disease.

Wound is a sore on the skin or a mucous membrane, accompanied by the disintegration of tissue. Wound can result in complete loss of the epidermis and often portions of the dermis and even subcutaneous fat. A wound that appears on the skin is often visible as an inflamed tissue with an area of reddened skin. Wound can also be caused due to a lack of mobility, which
causes prolonged pressure on the tissues. This stress in the blood circulation is transformed to a skin wound, commonly known as bedsores or decubitus ulcers.9

Patients may feel pain on the skin around the wound, and fluid may ooze from the wound. In some cases wound can bleed and, rarely patients experience fever. Ulcers develop in stages. In stage 1 the skin is red with soft underlying tissue.10 Chronic ulcer symptoms usually include increasing pain, friable granulation tissue, foul odour, and wound breakdown instead of healing (Figure 1).10 Ulcers may also appear on the cheeks, soft palate, the tongue, and on the inside of the lower lip. These ulcers usually last from 7 to 14 days and can be painful.11

Etiology

In addition to poor circulation, neuropathy, and difficulty in moving, there are factors that contribute to chronic wounds include systemic illnesses, age, and repeated trauma. Comorbid ailments that may contribute to the formation of chronic wounds include vasculitis, immune suppression, pyoderma gangrenosum, and diseases that cause ischemia.2 Immune suppression can be caused by illnesses or medical drugs used over a long period, for example steroids.2 Emotional stress can also negatively affect the healing of a wound, possibly by raising blood pressure and levels of cortisol, which lowers immunity.5 Another factor that may contribute to chronic wounds is old age.6

Comorbid factors that can lead to ischemia are especially likely to contribute to chronic wounds. Such factors include chronic fibrosis, edema, sickle cell disease, and peripheral artery disease such as by atherosclerosis.12 Repeated physical trauma plays a role in chronic wound formation by continually initiating the inflammatory cascade. The wounds from which ulcers arise can be caused by a wide variety of factors, but the main cause is impaired blood circulation. Especially, chronic wounds and ulcers are caused by poor circulation, either through cardiovascular issues or external pressure from a bed or a wheelchair.13 Other causes producing skin ulcers include bacterial or viral infections, fungal infections and cancers. Blood disorders and chronic wounds can result in skin ulcers as well.14

Pathophysiology

Chronic wounds may affect only the epidermis and dermis, or they may affect tissues all the way to the fascia.7

Classification of wounds: Wounds are classified by ‘stage’

Stage 1 wounds are characterized by redness or discoloration, warmth, and swelling or hardness. Stage 2 wounds partially penetrate the skin. Stage 3 describes full-thickness wounds that do not penetrate the tough white membrane (fascia) separating the skin and fat from the deeper tissues. Stage 4 wounds involve damage to muscle or bone and undermining of adjacent tissue. They may also involve the sinus tracts.7

The stages of wound healing

All wounds heal in three stages: (i) Inflammatory Stage, occurring during the first
few days. The wounded area attempts to restore its normal state by constricting blood vessels to control bleeding. Platelets and thromboplastin make a clot. Inflammation (redness, heat, swelling) also occurs and is a visible indicator of the immune response. White blood cells clean the wound of debris and bacteria. (ii) Proliferative Stage, lasting about 3 weeks (or longer, depending on the severity of the wound). Granulation occurs, which means that special cells called fibroblasts make collagen to fill in the wound. New blood vessels form. The wound gradually contracts and is covered by a layer of skin, and (iii) Maturation and Remodeling Stage, lasting up to 2 years. New collagen forms, changing the shape of the wound and increasing strength of tissue in the area. Scar tissue, however, is only about 80% as strong as the original tissue.

**Wound Infection**

Infection of a wound with a large number of bacteria, a process known as colonization, will slow the healing process. The difference between contamination and colonization is the concentration of bacteria. Anaerobic bacteria such as *Bacteroides*, *Clostridium* and *Streptococcus* may be active at deeper levels of the dermis, insulated from the healing influence of oxygen. Anaerobic bacteria are responsible for many devastating infections resulting in gangrene. Aerobic bacteria are more closely identified with superficial epidermal layers but may also be involved in infective processes and include *Staphylococcus epidermis*, *Corynebacteria*, and *Propionibacteria*.14

**Excessive use of antibiotics in non-healing ulcers**

Although approximately 4 million cases of non-healing ulcers are diagnosed annually in Europe, non-healing ulcers have been considered a negligible problem in society.14 Patients with non-healing ulcers is an excess usage of antibiotics. As early as 1998, it was reported that 60.1% of all ulcer patients were treated with at least one antibiotic within a six-month period.15

**Microbiology of chronic ulcers**

Conditions such as chronic venous insufficiency, arterial insufficiency, and pressure over time, can lead to the reduced reparation capacity of skin injuries, which can lead to non-healing ulcers.

A non-healing ulcer, however, should not be regarded as a disease, but rather as a symptom of an underlying state. Bacteria will colonize within the ulcer if the protective barrier of the skin is broken. Therefore, the appearance of a chronic ulcer depends on several factors (Table 1). These factors also contribute to the development of infections in the ulcer.16 Bacteria are always found in chronic ulcers. There are often multiple types of bacteria observed within a single ulcer.

For example, the flora usually found in cases of venous ulcers of the legs include *Staphylococcus aureus* (90.5%), *Enterococcus faecalis* (71.7%), and *Pseudomonas aeruginosa* (52.2%).17

The bacterial flora found in a non-healing ulcer change as the ulcer ages. *Staphylococci* and *Streptococci* bacteria are normally found in new ulcers, while gram negative mixed flora are often found in older ulcers. In addition, different types of ulcers are influenced by different types of bacteria. For example, a clinical infection will develop in 60% of diabetic foot ulcers but only 20% of venous leg ulcers that are colonized by *Staphylococcus aureus*.18 Between 1.6 and

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**Table 1: Factors affecting appearance of a chronic ulcer**

| Systemic factors                                                                 | Local factors            |
|---------------------------------------------------------------------------------|-------------------------|
| Metabolic diseases, such as diabetes mellitus                                  | Size of the ulcer       |
| Systemic diseases, such as rheumatic diseases                                   | Age of the ulcer        |
| Other forms of chronic disease, such as HIV infection                           | Location of the ulcer    |
| Old age                                                                         | Local circulation       |
| Malnutrition / poor diet                                                        | Necrosis                |
| Alcohol / narcotics abuse                                                       | Suppuration and maceration|
| Medicines, such as steroids, oestrogens, and vitamin K antagonists               | Edema                   |
| Smoking                                                                         | Exposed bones or capsules|

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4.4 species of bacteria are found per ulcer by conventional culturing methods. However, molecular biological methods suggest that even more species of bacteria are present in the average ulcer.

The number of ulcers with anaerobic bacterial growth is estimated to be between 25% and 82%. The most common anaerobic bacterial species are Peptostreptococcus and Prevotella. Recent research has indicated that the presence of bacterial biofilm contributes to the development of chronic ulcers. Studies performed earlier have shown that biofilm is present in 60% of chronic ulcers but only 6% of acute ulcers. This supports the view that biofilm probably plays an important role in the formation of chronic ulcerations.

**Treatment and prevention**

**Wound Treatment:** Hyperbaric Oxygen Therapy is used to treat very serious wounds. The patient breathes 100% oxygen in a pressurized chamber for 90-120 minutes. The oxygen dissolves into the blood and is distributed throughout the body, providing extra oxygen to the cells attempting to heal the wound. Hyperbaric oxygen treatments have been found to increase the rate of collagen deposition, angiogenesis, and bacterial clearance. Another benefit is that, if the wound environment has more oxygen, certain types of bacteria that cause serious infections cannot grow. This method has been used for many years in treating difficult or complicated, non-healing wounds.

It is well recognized as a very effective treatment. Skin ulcers may take a very long time to heal. Treatment is typically to avoid the ulcer getting infected, remove any excess discharge, maintain a moist wound environment, control the edema, and ease pain caused by nerve and tissue damage. Topical antibiotics are normally used to prevent the ulcer getting infected, and the wound or ulcer is usually kept clear of dead tissue through surgical debridement. Whirlpool Therapy is used by physical therapists once or twice daily for about 20 minutes during the inflammatory stage of healing to enhance circulation and bring more oxygen into the wound area. The whirlpool also softens and loosens dead tissue and cleanses the wound.

Ultrasound treatment uses mechanical vibration delivered at a frequency above the range of human hearing. Physical therapists report that covering the wound area with a hydrogel film and applying ultrasound during the inflammatory and proliferative stages stimulates the cells involved in wound healing and also warms the tissue, enhancing healing by improving circulation. Electrical Stimulation mimics the body’s own bioelectric system that influences wound healing by attracting repair cells, changing the permeability of cell membranes, and therefore affecting secretions and orienting cell structures.

**Nutritional supplements**

Researchers have shown that certain nutrients such as Aloe vera and vitamin C play key roles in wound healing. The typical Western diet is deficient in these nutrients. Injury significantly increases the need for the amino acid arginine, which is essential for a variety of metabolic functions. Arginine stimulates the cell-mediated immune response and protects against bacterial challenges. The amino acid glutamine is an important substrate for rapidly proliferating cells, including lymphocytes (white blood cells). It is also the major amino acid lost during muscle protein catabolism in the initial response to injury. Aloe vera provides the micronutrients required for protein synthesis. Its many components work together to reduce inflammation and pain, promote healing, and stop infection. Aloe can be applied topically to wounds and taken internally for both skin wounds and gastrointestinal ulcers.

Curcumin is an extract of the spice turmeric, known to have antioxidant properties and other health benefits. In Indian medicine, curcumin is used to reduce inflammation and treat wounds and skin ulcers. Topical application of curcumin encourages wound remodeling via effects on transforming growth factor-beta (TGF-b). It also improves reepitheliazation (new skin formation) and migration of cells such as myofibroblasts, fibroblasts, and macrophages, necessary for healing at the wound site.

Bromelain is found in pineapple and contains a proteolytic enzyme with the ability to break down or dissolve proteins. This mechanism of action can be helpful in chronic wounds or wounds having too much scar tissue. According to the PDR for Nutritional Supplements, bromelain speeds up healing time after surgical procedures. A German physician first observed the role of copper in healing too.
system is adversely affected by even moderate degrees of zinc deficiency. Severe zinc deficiency depresses immune function. Zinc is required for the development and activation of T-lymphocytes, a kind of white blood cell that helps fight infection. Zinc can be used topically or orally to encourage wound healing and plays a well-documented role in wound healing.33

Vitamin C is crucial for the proper function of the enzyme procollagen hydroxylase which produces collagen, the primary constituent of the granulation tissue that heals a wound and the key component in blood vessel walls. A published review stated that vitamin C plays a variety of roles in the prevention and treatment of cancer, including stimulating the immune system and enhancing wound healing. Wound healing requires more vitamin C than diet alone can easily provide. It must be replenished daily because it is water-soluble; any excess is excreted rather than stored.34 Vitamin B5 (Pantothenic Acid) improves healing by encouraging the migration of cells into the wounded area.35 Vitamin A is important for tissue synthesis and enhances resistance to infection.36 Vitamin B-Complex are needed for cell proliferation and for the replacement and maturation of red blood cells lost through bleeding.37

Different types of discharges from ulcer are

Serous, usually is seen in healing ulcers, while purulent is seen in infected ulcer. Yellow creamy discharge is observed in Staphylococcal infection; bloody opalescent discharge in Streptococcal infection, while greenish discharge is seen in Pseudomonas ulcers.4

Ulcer grading: Wagner’s grading of ulcer follow

Grade Description: 0: Pre-ulcerative lesion or healed ulcer, 1: Superficial ulcer, 2: Ulcer deeper to subcutaneous tissue exposing soft tissue or ulcer bone, 3: Abscess Formation Underneath, osteomyelitis, 4: Gangrene of part of tissues, limb or foot, and 5: Gangrene of entire one area or foot. Bloody (sanguineous) is usually seen in malignant ulcers and in healing ulcers with healthy granulation tissue, Seropurulent, Serosanguinous, Serous with sulphur granules, seen in actinomycosis, Yellowish, as seen in tuberculous ulcer.4

Complications

With any type of wound— even seemingly minor injuries— there is always danger of rapid multiplication of bacteria. The elderly and persons with reduced immunity are at great risk for wound-related infections. Once bacteria escape from the primary location of a wound, they enter the blood. This condition is commonly called blood poisoning, septicemia, sepsis, or septic shock. Sepsis is always a serious, life-threatening condition, with 56% mortality. In the United States, sepsis occurs annually in some three cases per 1,000 population. In sepsis and septic shock, circulation is reduced; blood pressure is markedly reduced, causing vital organs to receive reduced blood supply; heart, kidney, and liver functions are reduced or show signs of shutdown (multiple organ failure); and abnormal bleeding can develop. Symptoms of septicemia and septic shock are sudden onset of illness, high fever, chills, rapid breathing, headache, and altered mental state.38,39 Zaidi (2016) reported an option in treatment of poorly healing wounds with hirudotherapy (Figure 2-4).40

Fig. 2: Application of leech on chronic non-healing ulcer.

Fig. 3: Partially healed chronic ulcer.
DISCUSSION

Clinicians’ understanding of and ability to achieve wound healing has increased significantly over the past few years, particularly as a result of advances in molecular biology such as the use of growth factors, the ability to grow cells in vitro and the development of bioengineered tissue.\textsuperscript{41-43} Knowledge of scarring has also increased fundamentally.\textsuperscript{44-47} Some promising results have been obtained using epidermal growth factor,\textsuperscript{48} and keratinocyte growth factor-2\textsuperscript{49} for venous ulcers, and fibroblast growth factor\textsuperscript{50} and platelet-derived growth factor (PDGF) for pressure ulcers.\textsuperscript{51,52}

It is also possible that closer attention should have been paid to appropriately preparing the chronic wound before treatment with the growth factor being tested. Notably, there is evidence that the aggressive approach to surgical debridement taken in the initial PDGF trial for diabetic neuropathic ulcers seems to have worked synergistically with the application of the growth factor.\textsuperscript{53} Bioengineered skin products or skin equivalents have become available for the treatment of acute and chronic wounds as well as burns. Since the initial use of keratinocyte sheets,\textsuperscript{21,22,41} several more complex constructs have been developed and tested in human wounds. Skin equivalents may contain living cells, such as fibroblasts or keratinocytes, or both,\textsuperscript{2,41-43} while others are made of acellular materials or extracts of living cells.\textsuperscript{53}

Saline-soaked gauze and off-loading have been accepted by the Food and Drug Administration as the control. Bioengineered skin may work by delivering living cells which are known as a “smart material” because they are capable of adapting to their environment. There is evidence that some of these living constructs are able to release growth factors and cytokines.\textsuperscript{54,55} The technology to introduce certain genes into wounds by a variety of physical means or biological vectors, including viruses, has existed for some time. Work with gene therapy in relation to wounds has been done in experimental animal models.\textsuperscript{56}

There are promising indications that certain approaches may work in humans. For example, the introduction of naked plasmid DNA encoding the gene for vascular endothelial growth factor (VEGF) has been reported to enhance healing and angiogenesis in selected patients with ulcers resulting from arterial insufficiency.\textsuperscript{56}

CONFLICT OF INTEREST

The authors declare no conflict of interest.

REFERENCES

1 Mustoe T. Dermal ulcer healing: Advances in understanding. Tissue repair and ulcer/wound healing: molecular mechanisms, therapeutic targets and future directions. Paris, France: EUROCONFERENCES. Archived from the original (PDF) on October 27, 2005 (March 17–18, 2005).
2 Snyder, Robert J. Treatment of nonhealing ulcers with allografts. \textit{Clin Dermatol} 2005;\textit{23}:388–95.
3 Taylor JE, Laity PR, Hicks J, Wong SS, Norris K, Khunkhamchoo P, Johnson AF, Cameron RE. Extent of iron pick-up in deferoxamine-coupled polyurethane materials for therapy of chronic wounds. \textit{Biomaterials} 2005;\textit{26}:6024–33.
4 Krasner D. Painful venous ulcers: Themes and stories about living with the pain and

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\caption{Completely healed chronic non-healing ulcer after heridotherapy.}
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suffering. *J Wound Ostomy Continence Nurs* 1998;25:158–68.

5. Augustin M, Maier K. Psychosomatic Aspects of Chronic Wounds. *Dermatol Psychosomatics* 2003;4:5–13.

6. Mustoe T. Understanding chronic wounds: A unifying hypothesis on their pathogenesis and implications for therapy. *Am J Surg* 2004;187:S65.

7. Crovetti GI, Martinelli G, Issi M, Barone M, Guizzardi M, Campanati B, Moroni M, Carabelli A. Platelet gel for healing cutaneous chronic wounds. *Transfus Apher Sci* 2004;30:145-51.

8. Walsche C. Living with a venous leg ulcer: A descriptive study of patients’ experiences. *J Adv Nurs* 1995;22:1092-100.

9. Kumar V, Fausto N, Abbas A. Robbins & Cotran Pathologic Basis of Disease (7th ed.). Saunders. 2004; P. 1230.

10. Symptoms”. Retrieved 2010-06-16

11. Bhat S. Srb’s Manual of Surgery. 4e. Jaypee Brothers Medical Pub. 2013; p. 17.

12. Bhat S. Srb’s Manual of Surgery. 4e. Jaypee Brothers Medical Pub. 2013; p. 17.

13. Wong JK, Amin K, Dumville JC. The resource impact of wounds on healthcare providers in Europe. *J Wound Care* 2009;18:154.

14. Tammelin A, Lindholm C, Hambraeus A. Chronic ulcers and antibiotic treatment. *J Wound Care* 1998;7:435-7.

15. Edwards R, Harding KG. Bacteria and wound healing. *Curr Opin Infect Dis* 2004;17:91-6

16. Gjødsbol K, Christensen JJ, Karlsmark T, Jørgensen B, Klein BM, Krogfelt KA. Multiple bacterial species reside in chronic wounds: a longitudinal study. *Int Wound J* 2006;3:225-31.

17. Schmidt K, Debus ES, Jessberger S, Ziegler U, Thiede A. Bacterial population of chronic crural ulcers: is there a difference between the diabetic, the venous and the arterial ulcer. *VASA* 2000;29:62-70.

18. Bowler PG, Davies BJ. The microbiology of infected and noninfected leg ulcers. *Int J Dermatol* 1999;38:573-578.

19. Trengrove NJ, Stacey MC, McGechie, Mata S. Qualitative bacteriology and leg ulcer healing. *J Wound Care* 1996;5:277-80.

20. Parvez N, Dutta P, Ray P, Shah VN, Prakash M, Khandelwal N, Kaman L, Bhansali A. Microbial profile and utility of soft tissue, pus, and bone cultures in diagnosing diabetic foot infections. *Diabetes Technol Ther* 2012;14:669-74.

21. James GA, Swoffeger E, Wolcott R, Pulcini Ed, Secor P, Sestrich J, Costerton JW, Stewart PS. Biofilms in chronic wounds. *Wound Repair Regen* 2008;16:37-44

22. Pani U. Hyperbaric oxygen therapy as adjuvant in stump surgical wound healing. *G Ital Med Lav Ergon* 2015;37(Suppl):24-6.

23. Alhajj M, Nelson NG, McKenzie LB. Hot tub, whirlpool, and spa-related injuries in the U.S., 1990-2007. *Am J Prev Med* 2009;37:531-6.

24. Xia XP, Chen HL, Zhou B. Ultrasonography for meniscal injuries in knee joint: a systematic review and meta-analysis. *J Sports Med Phys Fitness* 2016;56:1179-1187.

25. Akhoondinasab MR, Akhoondinasab M, Saberi M. Comparison of Healing Effect of Aloe Vera Extract and Silver Sulfadiazine in Burn Injuries in Experimental Rat Model. *World J Plast Surg* 2014;3(1):29-34.

26. Vaxman F, Chalkiadakis G, Olender S, James GA, Swogger E, Wolcott R, Pulcini Ed, Ed, Maldonado H, Aprahamian M, Bruch JF, Wittmann T, Volkmar P, Grenier JF. Improvement in the healing of colonic anastomoses by vitamin B5 and C supplements. Experimental study in the rabbit. *Ann Chir* 1990;44:512-20.

27. Gurbuz AT, Kunzelman J, Ratzer EE. Supplemental dietary arginine accelerates intestinal mucosal regeneration and enhances bacterial clearance following radiation enteritis in rats. *J Surg Res* 1998;74:149-54.

28. Mehрабانی D, Farjam M, Geramizadeh B, Tanideh N, Amini M, Panjehshahin MR. The Healing Effect of Curcumin on Burn Wounds in Rat. *World J Plast Surg* 2015;4:29-35.

29. Rabbani Haghighi N, Naghsh N, Mehrabani D. The comparison of pretreatment effects of boiled coffee and curcuruma longa on serum albumin as a liver indicator in male rats injected with thioacetamide. *Fasa Univ Med Sci* 2014;4:58-66.

30. Aichele K, Bubel M, Deubel G, Pohlemann T, Oberringer M. Bromelain down-regulates myofibroblast differentiation in an in vitro wound healing assay. *Naunyn Schmiedebergs Arch Pharmacol* 2013;386:853-63.
32 Das A, Sudhahar V, Chen GF, Kim HW, Youn SW, Finney L, Vogt S, Yang J, Kweon J, Surenkhuu B, Ushio-Fukai M, Fukai T. Endothelial Antioxidant-1: a Key Mediator of Copper-dependent Wound Healing in vivo. Sci Rep 2016;6:33783.
33 Aydoğdu O, Tuncel U, Gümiş M, Kurt A, Oztürk N, Aksakal 1A, Çelik UR, Güzel N, Erkorkmaz U. Zinc-coated Foam With Negative Pressure Wound Therapy in the Treatment of Challenging Wounds: A New Alternative Interface Material. Wounds 2016 Aug 15. pii: WNDS20160815-2. [Epub ahead of print]
34 Mohammed BM, Fisher BJ, Kraskauskas D, Ward S, Wayne JS, Brophy DF, Fowler AA 3rd, Yager DR, Natarajan R. Vitamin C promotes wound healing through novel pleiotropic mechanisms. Int Wound J 2016;13:572-84.
35 Vaxman F, Olender S, Lambert A, Nisand G, Martinelli-Kläy CP, Lunardi LO, Martinelli C. Modulation of MCP-1, TGF-β1, and α-SMA Expressions in Granulation Tissue of Cutaneous Wounds Treated with Local Vitamin B Complex: An Experimental Study. Dermatopathology (Basel) 2014;1:98-107.
36 Egger SF, Huber-Spitzy V, Alzner E, Scholda C, Vecsei VP. Corneal wound healing after superficial foreign body injury: vitamin A and dexamethasone versus a calf blood extract. A randomized double-blind study. Ophthalmologica 1999;213:246-9.
37 Martinelli-Kläy CP, Lunardi LO, Martinelli CR, Lombardi T, Soares EG, Martinelli C. Modulation of MCP-1, TGF-β1, and α-SMA Expressions in Granulation Tissue of Cutaneous Wounds Treated with Local Vitamin B Complex: An Experimental Study. Dermatopathology (Basel) 2014;1:98-107.
38 Manafi A, Kohanteb I, Mehrabani D, Japoni A, Amini M, Naghmach M, Zaghie AH, Khalili N. Active immunization using exotoxin A confers protection against Pseudomonas aeruginosa infection in a mouse burn model. BMC J Microbiol 2009;9:19-23.
39 Tanideh N, Rokhsari P, Mehrabani D, Mohammadi Samani S, Sabet Sarvestani F, Ashraf MJ, Koohi Hosseinabadi O, Shamsian Sh, Ahmadi N. The Healing Effect of Licorice on Pseudomonas aeruginosa Infected Burn Wounds in Experimental Rat Model. World J Plast Surg 2014;3:99-106.
40 Zaidi SM. Unani treatment and leech therapy saved the diabetic foot of a patient from amputation. Int Wound J 2016;13:263-4.
41 Bell E, Ehrlich HP, Buttle DJ, Nakatsuji T. Living tissue formed in vitro and accepted as skin-equivalent tissue of full thickness. Science 1981;211:1052-54.
42 Boyce ST. Design principles for composition and performance of cultured skin substitutes. Burns 2001;27:523-33.
43 European Wound Management Association. Position Document: Wound Bed Preparation in Practice. London: MEP Ltd, 2004.
44 Longaker MT, Chiu ES, Adzick NS, Stern M, Harrison MR, Stern R. Studies in fatal wound healing. V. A prolonged presence of hyaluronic acid characterizes fetal wound fluid. Ann Surg 1991;213:292-96.
45 Longaker MT, Whitby DJ, Ferguson MW, Lorenz HP, Harrison MR, Adzick NS. Adult skin wounds in the fetal environment heal with scar formation. Ann Surg 1994;219:65-72.
46 Mackool RJ, Gittes GK, Longaker MT. Scarless healing. The fetal wound. Clin Plast Surg 1998;25:357-65.
47 Mast BA, Diegelmann RF, Krummel TM, Cohen IK. Scarless wound healing in the mammalian fetus. Surg Gynecol Obstet 1992;174:441-51.
48 Falanga V, Eaglstein WH, Bucalo B, Katz MH, Harris B, Carson P. Topical use of human recombinant epidermal growth factor (h-EGF) in venous ulcers. J Dermatol Surg Oncol 1992;18:604-6.
49 Robson MC, Phillips TJ, Falanga V, Odenheimer DJ, Parish LC, Jensen JL, Steed DL. Randomized trial of topically applied repifermin (recombinant human keratinocyte growth factor-2) to accelerate wound healing in venous ulcers. Wound Repair Regen 2001;9:347-52.
50 Robson MC, Phillips LG, Lawrence WT, Bishop JB, Youngerman JS, Hayward PG, Broemeling LD, Heggers JP. The safety and effect of topically applied recombinant basic fibroblast growth factor on the healing of chronic pressure sores. Ann Surg 1992;216:401-6.
51 Robson MC, Phillips LG, Thomason A, Robson LE, Pierce GF. Platelet-derived growth factor BB for the treatment of chronic pressure ulcers. Lancet 1992;339:23-25.
52 Pierce GF, Tarpley JE, Allman RM, Goode
PS, Serdar CM, Morris B, Mustoe TA, Vande Berg J. Tissue repair processes in healing chronic pressure ulcers treated with recombinant platelet-derived growth factor BB. *Am J Pathol* 1994;145:1399-410.

53 Veves A, Falanga V, Armstrong DG, Sabolinski ML, Apligraf Diabetic Foot Ulcer Study. Graftskin, a human skin equivalent, is effective in the management of noninfected neuropathic diabetic foot ulcers: a prospective randomized multicenter clinical trial. *Diabetes Care* 2001;24:290-95.

54 Mansbridge J, Liu K, Patch R, Symons K, Pinney E. Three-dimensional fibroblast culture implant for the treatment of diabetic foot ulcers: metabolic activity and therapeutic range. *Tissue Eng* 1998;4:403-14.

55 Falanga V, Isaacs C, Paquette D, Downing G, Kouttab N, Butmarc J, Badiavas E, Hardin-Young J. Wounding of bioengineered skin: cellular and molecular aspects after injury. *J Invest Dermatol* 2002;119:653-60.

56 Isner JM, Baumgartner I, Rauh G, Schainfeld R, Blair R, Manor O, Razvi S, Symes JF. Treatment of thromboangiitis obliterans (Buerger’s disease) by intramuscular gene transfer of vascular endothelial growth factor: preliminary clinical results. *J Vasc Surg* 1998;28:964-73.

57 Quesenberry PJ, Colvin GA, Lambert JF, Frimberger AE, Dooner MS, McAuliffe CI, Miller C, Becker P, Badiavas E, Falanga VJ, Elfenbein G, Lum LG. The new stem cell biology. *Trans Am Clin Climatol Assoc* 2002;113:182-206.