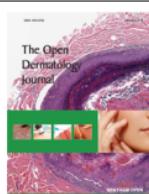


The Benefits Of Occlusive Dressings In Wound Healing

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REVIEW ARTICLE

The Benefits Of Occlusive Dressings In Wound Healing

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Abstract: There are several types of wounds with their own healing properties. The latest innovation in wound management by using occlusive dressings can prevent infections, improve healing time and patient's comfort. Occlusive dressings are often used as an immediate wound hygiene control and also prevent blood loss until debridement is performed. They are used to protect wounds and surrounding tissue from pathogens and other harmful materials. A good cover depends on the condition around the wound, the person's skills, and the injury's nature. In this article, we provide an insight into the types of polymer materials used clinically in wound dressing and underlying mechanisms between the biomaterial dressings and the body tissue.

Keywords: Occlusive dressings, Moist wound environment, Wound healing, Chronic wounds, Acute wounds, Epidermis.

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1. INTRODUCTION

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In general, wounds can be classified into acute and chronic wounds. Acute wounds mostly tend to heal completely within 8–12 weeks with minimal scarring. Chronic wounds tend to reoccur and 43 have a prolonged healing time that extends beyond 12 weeks. Wounds can also be classified according to 511 layers and affected areas. Superficial wounds involve only the epidermal skin surface. Partial thickness wounds involve the epidermis, deeper dermal layers, blood vessels, sweat glands, and hair follicles. Full thickness wounds are caused by injuries that extend into the subcutaneous fat or deeper tissue along with the epidermis and dermal layers [1, 2].

In 1958, it was the first time known that moist environment for the wound could improve wound healing, when it was found that blisters healed faster if left in their original form [3]. Previously, honey paste, plant fiber, and animal fat were commonly used. Gauzes, lint, and cotton wool began to be used as wound dressings a few years later [4]. It was found in several studies that a polyethylene film made partial-thickness wounds of porcine heal approximately two times faster than the wound left open to the air. After that, many studies agreed about the benefits of moist wound environment for acute wound healing [3]. This has been found in acute but not chronic wounds because no animal model exists for chronic wounds. Due to excessive wound drainage from the traditional dressings, it becomes moistened and tends to become adherent

to the wound making it painful when removing. These traditional dressings fail to provide moist environment to the wound so they have been replaced by modern dressings with more advanced formulations [5]. Moist condition makes wound healing faster, relieves pain, prevents infection and contamination than acute wound treated with other methods [3, 6]. To prevent infections, occlusive dressings need to be changed regularly. Occlusive dressings are mostly produced in the form of a flat shape sheet. Some of them can be cut according to the size of the wound, but some of them also need secondary dressings to completely seal the wound [7].

50 2. WOUND HEALING

Wound healing is a complex process which is influenced by various mechanisms involving the coordinated interaction of blood cells, proteins, proteases, growth factors, and extracellular matrix components. Wound healing process can be divided into four phases (1) the coagulation and haemostasis phase; (2) inflammatory phase; (3) proliferative phase; and (4) maturation phase [8].

Hemostasis is initiated during the exposure of collagen during wound formation that activates the intrinsic and extrinsic clotting cascade. Injury to tissue causes a release of thromboxane A2 and prostaglandin 2-alpha which leads to vasoconstriction. Extravasation of blood constituents assists in the formation of the blood clot. All of this process helps to limit hemorrhage and provides an initial extracellular matrix for cell migration.

After hemostasis is achieved, inflammatory phase begins. Capillary vasodilatation and leakage result secondary to local

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histamine release by the activated complement cascade. The increased blood flow and altered vascular permeability allow for the migration of inflammatory cells to the wound bed. Complement, neutrophil and macrophage play an important role in stimulating inflammatory cells, bacterial lysis and debris scavenging. In addition to direct phagocytosis of bacteria and foreign materials, macrophages secrete various enzymes and growth factors.

The proliferative phase is marked by epithelialization, angiogenesis, granulation tissue formation, and collagen deposition. Epithelialization occurs within hours after injury in wound healing and the normal layers of epidermis are restored typically in 3 days. The proliferative phase ends with granulation tissue formation.

The maturational phase is characterized by transition from the granulation tissue to scar formation. Around two weeks after injury, the wound undergoes contraction which results in a smaller amount of apparent scar tissue. Collagen deposition by fibroblasts continues for a prolonged period with a net increase in collagen deposition reaching after three weeks from tissue injury. The entire process is a dynamic continuum dictated by numerous growth factors and cells with an overlap of each of the three phases of wound healing to provide continued remodeling. The human wound has its maximal strength at one year [4, 9].

Promotion of these phases largely depends on the wound type, its associated pathological conditions and also the type of dressing material to achieve faster healing. Several factors can impair the whole process of wound healing [39]. In general, these factors are classified as local and systemic. Local factors that affect wound healing are desiccation, infection, maceration, necrosis, oxygenation, pressure, trauma, and edema. (Checklist wound healing). Systemic factors such as aging, hormones, stress, and systemic diseases are essential in determining individual wound healing process [8, 10 - 14].

3. BASICS OF OCCLUSIVE DRESSINGS

Based on the wound type, suitable dressing material must be used. Dressing selection should be based on its ability to a) provide or maintain moist environment, b) enhance epidermal migration, c) promote angiogenesis and connective tissue synthesis, d) allow gas exchange between wounded tissue and environment, e) maintain appropriate tissue temperature to improve the blood flow to the wound bed and enhance epidermal migration, f) provide protection against bacterial infection, g) should be non-adherent to the wound and easy to remove after healing, h) must provide debridement action to enhance leucocytes migration and support the accumulation of enzyme, and i) must be sterile, non-toxic and non-allergic [15].

Modern wound dressings are usually based on synthetic polymers and are classified as passive, interactive and bioactive products. Passive products are non-occlusive, such as gauze and tulle dressings, used to cover the wound to restore its function underneath. Interactive dressings are semi-occlusive or occlusive, available in the form of films, foam, hydrogel and hydrocolloids. These dressings act as a barrier against penetration of bacteria to the wound environment [16 - 19].

Occlusive dressings increase about 40% of the process of

re-epithelialization of partial thickness wound [4, 20]. Studies where occlusive dressing leads to earlier epithelialization during wound healing have been proposed [20]. When the wound is closed with dressing, they are continuously exposed to proteinases, chemotactic, complement [16], growth factors, which is lost in the wound exposed. These dressings help in faster re-epithelialization, collagen synthesis, promote angiogenesis by creating hypoxia to the wound bed and decrease wound bed pH which leads to a decrease in the wound infection [21]. An increase in oxygen tension under occlusion has also been suggested as a beneficial effect of these dressings; however, hydrocolloids, which have healing rates as good as or better than those of polymer films, are actually oxygen impermeable, whereas the films are oxygen permeable.

Inflammatory phase and angiogenesis are enhanced in moist wound environment when compared to dry wound environment. Based on Winter *et al.* studies, dry wound tends to heal slowly with poor cosmesis compared with wound in a moist environment due to impaired reepithelialisation and scab formation [16, 19]. Fig. (1) describes the basic mechanism of occlusive dressings.

The number of endothelial and fibroblast cells increases in moist wound conditions as compared to wounds that are kept dry. Inflammatory phase cells such as neutrophils and macrophages increase in moist wound environment which can reduce scab or debris formation. By preventing wound desiccation, occlusion maintains an electrical gradient between wounded and non-wounded skin, which may stimulate epidermal cell migration.

Fluid in the wound improves wound healing by interacting with regenerating epithelium, dermal, and granulation tissue (ie, fibroblasts and endothelial cells). An increase in oxygen tension in combination [53] with electrical stimulation has been shown to increase the expression of growth factor receptors on endothelial cells and fibroblast. Endothelial cells and fibroblast stimulated by platelet-derived growth factor increase collagen synthesis and promote angiogenesis. This mechanism plays an important role in preserving cosmesis. Inflammatory infiltrate develops from the dominant neutrophils to macrophages in mature wounds. Occlusion shortens the inflammatory healing phase and produces good cosmetic results. Fig. (2) describes the effects of occlusive dressings in wound healing process [4, 5].

It was thought that occlusion of wounds may lead to an increased risk of bacterial infection, but studies showed that wounds treated with dressings, which promote moist wound environment [7] are associated with a lower infection rate. Dermatitis has been reported after prolonged exposure to water, but such hydration induced changes to the epidermis can be reversed quickly if the underlying cause is removed. A wound dressing thus approaching ideal characteristics should conform to the site of the wound, offer alleviation of pain symptoms, promote faster wound-healing time and attempt to restore the patients' normal daily activities. Occlusive dressings protect the wound from both pathogenic invasion and further trauma. They act as barriers and prevent the entry of outside pathogens that might infect the wound and retard healing.

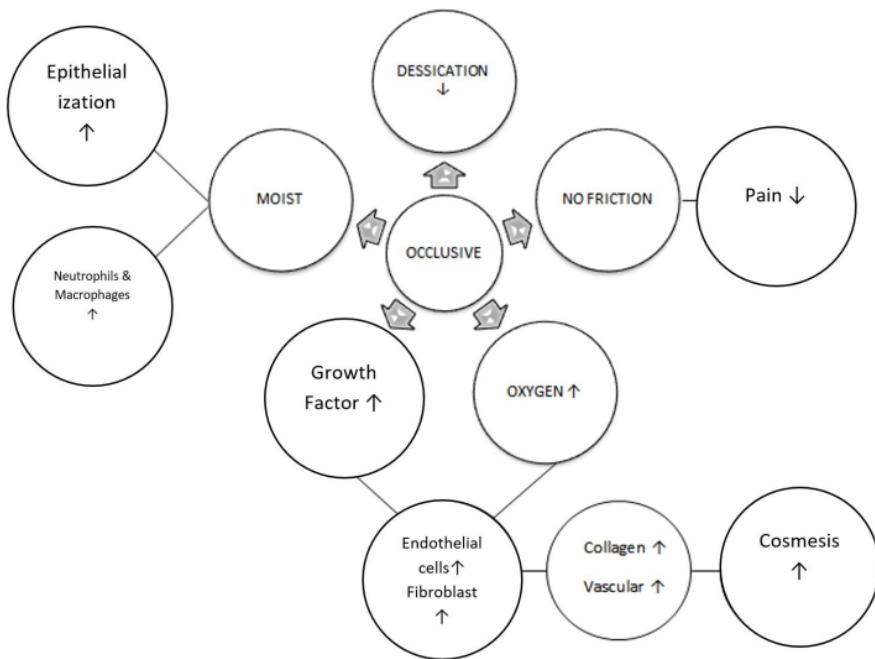


Fig. (1). Basic Mechanism of Occlusive Dressing.

1 ling. Since they conform to the body's contours, they are more likely to remain in place and offer protection from further trauma as well [4, 22 - 26].

23 4. INDICATIONS AND CONTRAINDICATIONS FOR OCCLUSIVE DRESSINGS

We summarized the **indications** and contraindications of occlusive dressings and also its beneficial effects.

Indications of occlusive dressings are [7]:

- Wound treatment. Occlusive dressings can be combined with antibiotics, gauze, sponge, hydrogels, and any other method to treat wounds
- Pressure and bleeding can often be immediately addressed
- Topical ointments. It gives pressure to the ointment which improves absorption into the wound and prevents evaporation.
- Partial evisceration. Occlusive dressings are applied to protect the bowel until surgery.
- Sucking chest wounds. Wound and the puncture are covered and treated.

Contraindications of occlusive dressings are [7]:

- Maceration of the skin, where the skin feels tender, moist, and looks whiter than before, is an occlusive dressing's long-term risk.

- Pathogenic germs that exist in the infected area. Benefits of occlusive dressings [6]:

- Improve patient's comfort
- Lower the risk of infection
- Speed up wound healing

Several studies show the benefits of occlusive dressing for treating herpes virus skin infection. Based on histological analysis, it is known that skin injury caused by herpes virus is similar to partial-thickness skin wounds. Based on the study by Keegan *et al.*, occlusive dressing relieved pain and hypersensitivity caused by herpes zoster infection which may lead as an option to rule out the need for oral analgesics. A blinded control study by Lin *et al.* showed that the use of plain Tegaderm™ provided better pain relief. Lee *et al.* also showed the benefits of occlusive dressing for the management of herpes zoster by improving wound healing, relieving pain, improving patient's comfort and eschar debridement [27 - 29].

5. TYPES OF OCCLUSIVE DRESSINGS

Occlusive dressings work by maintaining moist wound environment which can shield the wound surface by preventing dry environment and trauma. Dry environment can prevent new epidermal cells migration to the wound surface. Variant types are: 1) semipermeable films, 2) hydrogels, 3) hydrocolloids and 4) alginates. Examples, functions, and features of each follow [6].

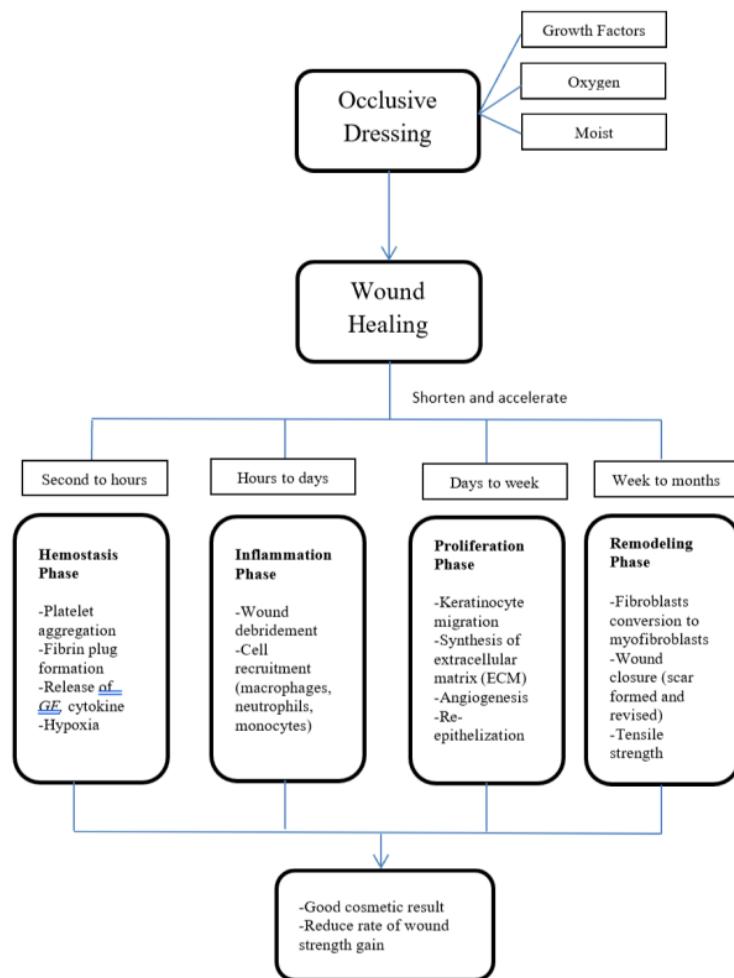


Fig. (2). Occlusive Dressing and Wound Healing Process.

5.1. Polymer Films

In 1960s, film dressings were introduced for wound management. These transparent semipermeable dressings are made of polyethylene or polyurethane with an adhesive coating on one side, are thin, highly elastomeric, elastic, permeable to moisture vapor and atmospheric gases but impermeable to liquid and bacteria [3]. Originally, these films are used for healing minor burns, superficial wounds, superficial decubitus ulcers, donor sites, blisters, abrasions, and cuts [6].

During the past years, wound dressing has been modernized which now mainly consists of synthetic polymers for wound management and can be classified as passive and interactive. Passive synthetic polymer dressings are non-occlusive, used for covering wound and help in restoring function under the polymer film. Examples of such passive synthetic polymers are gauze and tulle. Interactive synthetic

polymer dressings are occlusive or semi-occlusive which provide a barrier against bacterial penetration to the wound [30].

Polymer films trap exudates so the wound environment becomes moist [3, 4]. One main feature is that these films are impermeable to bacteria and liquid but are permeable to moisture vapor and air. The major drawback of polymer dressings is that exudates from wounds may accumulate underneath these dressings, which, due to increased pressure may cause a break in the environment maintained by the occlusive dressing [31, 32]. These products can be used for long-term bedridden patients acting as a shield to protect against skin friction in vulnerable areas, as a cover over topical products, and as intravenous dressings [3, 6].

5.2. Polymer Foams

Polyurethane foam film dressing is made of a thin microporous sheet of polyurethane foam that has been coated with a hydrophilic adhesive and bonded to a polyurethane film layer, which is good to avoid dehydration and bacterial infection. Foam film dressing has small pore size which increases the effectiveness of absorption or fluid retention. It also provides effectiveness in the prevention of bacterial infection and dehydration. Foam dressings attempt to rectify the lack of exudate absorbancy of occlusive dressing, without compromising on the moist environment needed for tissue repair [33 - 36]. Foams dressing does not cause maceration even if used for several days [4].

The major weakness of this type of dressing is that it needs secondary dressing like elastic bandage or a film to attach to the wound. Foam dressings are used for the wound that produced many exudates and deep wound or used under the compression bandage for chronic wound like venous ulcer [3, 4].

5.3. Hydrogel Dressings

Hydrogel contains about 96% water but does not dissolve in water and causes swelling if exposed to liquid solutions which can transmit moisture vapor. The high water content of hydrogels helps granulation tissues and epithelium in a moist environment. Hydrogel is made of polymers (polyethylene oxide, polyacrylamide, and polyvinylpyrrolidone) which are insoluble hydrophilic. Soft elastic property of hydrogels provides easy application and removal after the wound is healed without any damage. Their rate of absorbance is low, but high in capacity. For application, hydrogels need a secondary layer to attach to the wound, such as bandage or tape.³ Hydrogel has soothing and cooling effect up to 5°C and maintains reduction in the cutaneous wounds temperature for up to 6 hours (may be augmented by refrigeration before use) which can alleviate pain and inflammation.

Table 1. Wound Dressings Types and Purpose [1]

Type	Purpose Use For
Polymeric hydrogels	Ulcers Chemotherapy peels Laser resurfacing
Polymeric foam	Burns Chronic wounds Laser resurfacing wounds Mohs surgery and wounds
Polymeric hydrocolloides	Average thickness wounds Burns Chronic Ulcer
Polymeric alginates	Surgical wounds Thickness burns Chronic ulcer High exudate wounds

The main disadvantage for using hydrogel is that it is easy to dry out which reduces their effectiveness and needs to be changed after 2 days [6]. Hydrogel is also a bad bacterial barrier.³ Morgan has reported that except for infected and heavy drainage wounds, hydrogel dressings are suitable for all

four stages of wound healing. Difficulties in hydrogel dressings are that exudate accumulation leads to maceration and bacterial proliferation that produces foul smell in wounds. Besides, low mechanical strength of hydrogels makes it difficult to handle [36, 37]. Hydrogel is indicated for partial-thickness burns, dry chronic wounds, necrotic wounds, blisters and minor lacerations (Table 1) [3].

5.4. Hydrocolloid Dressings

Hydrocolloid is waterproof, semipermeable to vapor, opaque, occlusive, and absorbent. ¹³⁸ Hydrocolloid can be obtained synthetically or naturally [4]. It consists of two layers, inner colloidal layer and outer water-impermeable layer [3]. Hydrocolloids often use polysaccharides gel agents (sodium carboxymethyl cellulose), elastomeric compound, and are adhesive [4]. This type of dressing does not need secondary dressing and is impermeable to gases and moisture vapor [1]. Hydrocolloids are permeable to water vapor but impermeable to bacteria and also have the properties of debridement and absorb wound exudates [38]. They can melt so when they absorb exudate, but become viscous gel to stay in the wound when the dressings are removed. Hydrocolloids can be removed by normal saline [3].

The encapsulation properties of hydrocolloids create an opportunity to produce enhanced products with controlled drug (or any other appropriate substance) release due to the different type of hydrocolloid gels with different structural gelation. The amount of drug release can be controlled by optimization of the size of particles and the permeability of the gel membrane [39, 40].

Hydrocolloid dressings are ¹³ned on some wounds for prolonged periods (more than 1 week); this is useful in managing clean ulcers, but not when regular wound inspection is required. Thus, these dressings are probably more useful in preventing, rather than treating, infection within a wound [41]. The disadvantages of this dressing are the “gel and smell” phenomenon which may create an odorous residue when the exudate is mixed with adhesive when the dressing is removed.⁶ In medical applications, the current market for hydrocolloid dressings offers their use in healing diabetic foot ulcers, chronic wound management, burn, and are also recommended for paediatric wound care management, as they do not cause pain on removal [3, 4].

5.5. Alginates

¹¹ Alinate dressings are produced from the calcium and sodium salts of alginic acid which can be obtained from seaweed. Alinate dressings can be in the form of nonwoven mats, porous (foam) sheets or twisted staple fibers.³ Alinate works by ion exchange reaction. The dressings would swell when wound exudates or sodium ions in tissue ⁴⁵¹ are exchanged with alginates dressings of calcium ions. Alinate forms a gel in contact with the exudates. This property of alinate gels makes them very desirable as dressing for wounds that contain low to high exudates [42]. It would stay as gel for around one month if compared to hydrocolloids which degrade faster [3, 4].

Table 2. Occlusive Dressings Benefits¹

	Polymer Foams	Hydrogels	Polymer Films	Alginates	Hydrocolloids
Absorbent	+	+	-	+	±
Transparent	-	±	+	-	-
Suited for exudative wounds	+	±	-	+	-
Permeable to water vapor	+	+	+	+	-
Permeable to oxygen	+	+	+	+	-
Prevents bacterial entry	-	-	-	-	+
Adherent to nonwounded skin	-	-	+	-	+

One of the main reasons for using alginates in wound dressings is their haemostatic ability; therefore, alginate dressings can be used for wounds that are bleeding. The coagulation effects of zinc and calcium alginate dressings have been compared with non-alginate dressings. Zinc containing alginates have the best haemostatic ability. Alginate dressings have an important property of gel formation which helps in reducing the pain during removal and changing of the ³⁴age from the wound site. Alginate gels can also be used for the delivery of drugs having low molecular weight ³⁴ie partially oxidized alginate gels are known to be used for controlled and localized delivery of antineoplastic agents [43, 44].

Alginate dressings have applications in tissue regeneration and bio-engineering fields. Depending on composition, sodium alginate has been identified as a substrate for cell proliferation. This opens up new possibilities for tissue regeneration in skin scaffolds as well [45]. Even though some studies have reported that alginate inhibits keratinocytes migration, Thomas *et al.*, have reported that alginates accelerate healing process by activating macrophages to produce TNF- α which initiates inflammatory signals [46]. In dry condition, alginate dressing removal may damage the newly formed epithelium around the wound. Alginate dressings are suitable for moderate to heavy drainage wounds and not suggested for dry wound, third degree burn wound and severe wounds with exposed bone. Like polymer foam and hydrogel, alginate also needs a secondary dressing to be attached (Table 2) [3].

CONCLUSION

Wound dressings have experienced development to more active dressings which create moist wound environment. The ideal characteristics of an occlusive dressings are to maintain moist environment, shorten and accelerate wound healing process such as; enhance epidermal migration, promote angiogenesis and connective tissue synthesis, allow gas exchange, improve blood flow to the wound bed and enhance epidermal migration. Moreover, they provide protection against bacterial infection, are non-adherent to the wound providing debridement action, are sterile, non-toxic and non-allergic. Occlusive dressings are not suitable to treat wounds with impaired circulation such as burn ulcers, pressure ulcers, or other heavy trauma.

CONSENT FOR PUBLICATION

⁵ Written consent was obtained from the patient for the surgery and the publication of her data.

AVAILABILITY OF DATA AND MATERIALS

Not applicable.

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CONFLICT OF INTEREST

The authors declare no conflict of interest, financial or otherwise.

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REFERENCES

- [1] Bryant R, Nix D. Acute and chronic wounds. Amsterdam: Elsevier Health Sciences 2006.
- [2] Bolton L, van Rijswijk L. Wound dressings: Meeting clinical and biological needs. *Dermatol Nurs* 1991; **3**(3): 146-61. [PMID: 1828677]
- [3] Helfman T, Ovington L, Falanga V. Occlusive dressings and wound ⁸25. *J Clin Dermatol* 1994; **12**(1): 121-7. [http://dx.doi.org/10.1016/0738-081X(94)90262-3] [PMID: 8180934]
- [4] Mir M, Ali MN, Barakatullah A, *et al.* Synthetic polymeric biomaterials for wound healing: a review. *Prog Biomater* 2018; **7**(1): 1-21. [http://dx.doi.org/10.1007/s40204-018-0083-4] [PMID: 29446015]
- [5] Boateng JS, Matthews KH, Stevens HNE, Eccleston GM. Wound healing dressings and drug delivery systems: A review. *J Pharm Sci* 2008; **97**(8): 2892-93. [http://dx.doi.org/10.1002/jps.35] [PMID: 17963217]
- [6] Scot B, MS, ATC, PA-S. Wound Management: The Occlusive Dressing. *J Athl Train* 1995; **30**(2): 143-46.
- [7] WoundSource. What is an Occlusive Dressing?. 2016.
- [8] Dhivya S, Padma VV, Santhini E. Wound dressings - A review. *Biomedicine (Taipei)* 2015; **5**(4): 22. [http://dx.doi.org/10.7603/s40681-015-0022-9] [PMID: 26615539]
- [9] Simo H, Prakash S. Complements and the wound healing cascade: an ¹⁰133 ed review. *Plast Surg Int* 2013; **2013**(146764)146764 [http://dx.doi.org/10.1155/2013/146764] [PMID: 23984063]
- [10] Guo S, Diptro LA. Factors affecting wound healing. *J Dent Res* 2010; **89**(3): 219-29. [http://dx.doi.org/10.1177/0022034509359125] [PMID: 20139336]
- [11] Godbout JP, Glaser R. Stress-induced immune dysregulation: implications for wound healing, infectious disease and cancer. *J Neuroimmune Pharmacol* 2006; **1**(4): 421-7. [http://dx.doi.org/10.1007/s11481-006-9036-0] [PMID: 18040814]
- [12] Brem H, Tomic-Canic M. Cellular and molecular ²⁴33 of wound healing in diabetes. *J Clin Invest* 2007; **117**(5): 1219-22. [http://dx.doi.org/10.1172/JC132169] [PMID: 17476353]
- [13] Wilson JA, Clark JJ. Obesity: impediment to postsurgical ⁵²wound healing. *Adv Skin Wound Care* 2004; **17**(8): 426-35. [http://dx.doi.org/10.1097/00129334-200410000-00013] [PMID: 15492679]
- [14] Thomas Hess C. Checklist for factors affecting wound healing. *Adv Skin Wound Care* 2011; **24**(4): 192. [http://dx.doi.org/10.1097/01.ASW.0000396300.04173.ec] [PMID: 2142 2844]

- [15] Degreef HJ. How to heal a wound fast. *Dermatol Clin* 1998; 16(2): 365-75. [\[PMID: 9589210 \]](http://dx.doi.org/10.1016/S0733-8635(05)70019-X)

[16] Hunt TK, Hopf H, Hussain Z. Physiology of wound healing. *Adv Skin Wound Care* 2000; 13(2)(Suppl.): 6-11. [\[PMID: 11074996 \]](http://dx.doi.org/10.1074/j.jclinmed.2006.10.001)

[17] Rivera AE, Spencer JM. Clinical aspects of full-thickness wound healing. *Curr Dermatol* 2007; 25(1): 39-48. [\[PMID: 172766 \]](http://dx.doi.org/10.1016/j.jclinmed.2006.10.001)

[18] Streckler-McGraw MK, Jones TR, Baer DG. Soft tissue wounds and phases of healing. *Emerg Med Clin North Am* 2007; 25(1): 1-22. [\[PMID: 17400070 \]](http://dx.doi.org/10.1016/j.emc.2006.12.002)

[19] Winter GD. Effect of air exposure and occlusion on experimental human skin wounds. *Nature* 1963; 200: 378-9. [\[PMID: 14087905 \]](http://dx.doi.org/10.1038/200378a0)

[20] Nayeri F. Occlusive bandaging of wounds with decreased circulation promotes growth of anaerobic bacteria and necrosis: case report. *BMC Infect Dis* 2016; 9: 394. [\[PMID: 27502024 \]](http://dx.doi.org/10.1186/s13104-016-2205-1)

[21] Samah S. Recent advances in topical wound care. *Indian J Plast Surg* 2017; 47(4S): 379-87. [\[PMID: 23162238 \]](http://dx.doi.org/10.4103/0970-0358.101321)

[22] Winter GD, Scales JT. Effect of air drying and dressings on the surface of a wound. *Nature* 1963; 197: 91-2. [\[PMID: 14001241 \]](http://dx.doi.org/10.1038/197091b0)

[23] Ousey K, Cutting KG, Rogers AA, Rippon MG. The importance of hydration in wound healing: reinvigorating the clinical perspective. *J Wound Care* 2016; 25(3): 122-130, 124-130. [\[PMID: 26947692 \]](http://dx.doi.org/10.12968/jowc.2016.25.3.122)

[24] Björklund S, Ruzgas T, Nowacka A, et al. Skin membrane electrical impedance properties under the influence of a varying water gradient. *Biophys J* 2013; 104(12): 2639-50. [\[PMID: 23790372 \]](http://dx.doi.org/10.1523/j.biophys.2013.05.008)

[25] Thomas S. 2008 The role of dressings in the treatment of moisture-related skin damage. *World Wide Wounds* [February 23]; [\[PMID: 16558325 \]](http://bit.ly/1wRQThs)

[26] Rheincker SB. Wound management: the occlusive dressing. *J Athl Train* 1995; 30(2): 143-6. [\[PMID: 16558325 \]](http://dx.doi.org/10.1123/jat.30.2.143)

[27] Keegan DA. Reducing pain in acute herpes zoster with 44% lanolin occlusive dressings: A case report. *J Med Case Reports* 2015; 9: 89. [\[PMID: 25907451 \]](http://dx.doi.org/10.1186/s13256-015-060-5)

[28] Lin PL, Fan SZ, Huang CH, et al. Analgesic effect of lidocaine patch 5% in the treatment of acute herpes zoster: A double-blind and vehicle-controlled study. *Reg Anesth Pain Med* 2008; 33(4): 320-5. [\[PMID: 18675742 \]](http://dx.doi.org/10.1097/0011550-200807000-00006) [\[PMID: 26615539 \]](http://dx.doi.org/10.7603/s04681-015-0022-9)

[29] Lee SK. Notes on Practice: Healing Shingles with Moist Occlusive Dressings. *Wound Management & Prevention* 2002; 40(3)

[30] Dhivya S, Padma VV, Santhini E. Wound dressings - A review. *BioMedicine (Taipei)* 2015; 5(4): 22. [\[PMID: 26615539 \]](http://dx.doi.org/10.7603/s04681-015-0022-9)

[31] Jenks M, Craig J, Green W, Hewitt N, Arber M, Sims A. *Telegderm CHG IV* securement dressing for central venous and arterial catheter insertion sites: A NICE medical technology guidance. *Appl Health Econ Health Policy* 2016; 14(2): 135-49. [\[PMID: 26458938 \]](http://dx.doi.org/10.1007/s40258-015-0202-5)

[32] Khil MS, Cha DI, Kim HY, Kim IS, Bhattacharai N. Electrospun nanofibrous polyurethane membrane as wound dressing. *J Biomed Mater Res Part B: Appl Biomater* 2003; 67(2): 675-9. [\[PMID: 14598393 \]](http://dx.doi.org/10.1002/jbmb.10058)

[33] Hinrichs WL, Lommen EJ, Wildevuur CR, Feijen J. Fabrication and characterization of an asymmetric poly(46) membrane for use as a wound dressing. *J Appl Biomater* 1992; 3(4): 287-303. [\[PMID: 10147998 \]](http://dx.doi.org/10.1002/jab.20048)

[34] Helfman T, Ovington L, Falanga V. Occlusive dressings and wound healing. *Clin Dermatol* 1994; 12(1): 121-7. [\[PMID: 8180934 \]](http://dx.doi.org/10.1016/0881-081X(94)90262-3)

[35] Zahedi P, Rezaee-O A, Ranaei-Statad S, Jafari S, Supaphol P. A review on wound dressings with an emphasis on electroporous nanofibrous polymeric bandages. *Polym Adv Technol* 2010; 21: 77-95. [\[PMID: 2048966 \]](http://dx.doi.org/10.1002/pat.2048)

[36] Johnson T. *Foam Composite*. US Patent 20048966, 2006.

[37] Martin L, Wilson CG, Koosha F, et al. The release of model macromolecules may be controlled by the hydrophobicity of palmitoyl glycol chitosan hydrogels. *J Control Release* 2002; 80(1-3): 87-100. [\[PMID: 1194390 \]](http://dx.doi.org/10.1016/S0168-3659(02)00005-6)

[38] Thomas S, Loveless PA. A comparative study of twelve hydrocolloid dressings. *World Wide Wounds* 1997; 1: 1-12.

[39] Bramhill J, Ross S, Ross G. Bioactive nanocomposites for tissue repair and regeneration: A review. *Int J Environ Res Public Health* 2017; 14(1): 66. [\[PMID: 28085054 \]](http://dx.doi.org/10.3390/ijerph1401006)

[40] Das S, Baker AB. Biomaterials and nanotherapeutics for enhancing skin wound healing. *Front Bioeng Biotechnol* 2016; 4: 82. [\[PMID: 27843895 \]](http://dx.doi.org/10.3389/fbioe.2016.00051)

[41] Frykberg RG, Banks J. Challenges in the treatment of chronic wounds. *Adv Wound Care (New Rochelle)* 2015; 4(9): 560-82. [\[PMID: 26339534 \]](http://dx.doi.org/10.1089/wound.2015.0635)

[42] Dabiri G, Damstetter E, Phillips T. Choosing a wound dressing based on common wound characteristics. *Adv Wound Care (New Rochelle)* 2016; 5(1): 32-41. [\[PMID: 26858913 \]](http://dx.doi.org/10.1089/wound.2014.0586)

[43] Segal HC, Hunt BJ, Gilding K. The effects of alginate and non-alginate wound dressings on blood coagulation and platelet activation. *J Biomater Appl* 1998; 12(3): 249-57. [\[PMID: 9493071 \]](http://dx.doi.org/10.1177/088532829801200305)

[44] Paul W, Sharma CP. Chitosan and alginate wound dressings. *Trends in Wound Artif Organs* 2004; 18(1): 18-23.

[45] Wang L, Shelton RM, Cooper PR, Lawson M, Triffitt JT, Barralet JE. Evaluation of sodium alginate for bone marrow cell tissue engineering. *Biomaterials* 2003; 24(20): 3475-81. [\[PMID: 12809776 \]](http://dx.doi.org/10.1016/S0101-4429(03)00167-4)

[46] Thomas A, Harding KG, Moore K. Alginates from wound dressings activate human macrophages to secrete tumour necrosis factor-alpha. *Biomaterials* 2000; 21(17): 1797-802. [\[PMID: 10905462 \]](http://dx.doi.org/10.1016/S0101-4429(00)00072-7)

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| | Publication | |
| 3 | www.magonlinelibrary.com | % 1 |
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| | Student Paper | |
| 7 | Submitted to Walla Walla College | % 1 |
| | Student Paper | |
| | Submitted to University of Leeds | |

9 Charles E. Argoff. "Introduction", Postgraduate Medicine, 2015

Publication

% 1

10 Aoki, Shigehisa, Toshiaki Takezawa, Satoshi Ikeda, Yutaka Narisawa, Ayumi Oshikata-Miyazaki, Syohei Miyauchi, Hiroshi Hirayama, Tomoya Sawaguchi, Tomoyuki Chimuro, and Shuji Toda. "A new cell-free bandage-type artificial skin for cutaneous wounds", Wound Repair and Regeneration, 2015.

Publication

% 1

11 Submitted to University of Huddersfield

Student Paper

% 1

12 Steven L. Percival. "Restoring balance: biofilms and wound dressings", Journal of Wound Care, 2018

Publication

% 1

13 cid.oxfordjournals.org

Internet Source

% 1

14 L.V. Arsenie, C. Pinse, A. Bethry, L. Valot, P. Verdie, B. Nottelet, G. Subra, V. Darcos, X. Garric. "Star-poly(lactide)-peptide hybrid networks as bioactive materials", European Polymer Journal, 2020

% 1

- 15 Li, M.. "Electrospinning polyaniline-contained gelatin nanofibers for tissue engineering applications", *Biomaterials*, 200605 % 1
Publication
-
- 16 Submitted to La Trobe University % 1
Student Paper
-
- 17 discovery.ucl.ac.uk % 1
Internet Source
-
- 18 Charernsriwilaiwat, Natthan, Theerasak Rojanarata, Tanasait Ngawhirunpat, and Praneet Opanasopit. "Electrospun chitosan/polyvinyl alcohol nanofibre mats for wound healing", *International Wound Journal*, 2012. % 1
Publication
-
- 19 www.woundsresearch.com % 1
Internet Source
-
- 20 www.tandfonline.com <% 1
Internet Source
-
- 21 Pieter R. Zwanenburg, Berend T. Tol, Miryam C. Obdeijn, Oren Lapid, Sarah L. Gans, Marja A. Boermeester. "Meta-analysis, Meta-regression, and GRADE Assessment of Randomized and Nonrandomized Studies of Incisional Negative Pressure Wound Therapy Versus Control <% 1

Dressings for the Prevention of Postoperative Wound Complications", Annals of Surgery, 2020

Publication

-
- 22 www.labome.org <% 1
Internet Source
-
- 23 www.woundsource.com <% 1
Internet Source
-
- 24 journalofethics.ama-assn.org <% 1
Internet Source
-
- 25 Submitted to University of Queensland <% 1
Student Paper
-
- 26 Stem Cell Biology and Regenerative Medicine, <% 1
2015.
Publication
-
- 27 Allan Lemos Maia, Esdras Marques Lins, José <% 1
Lamartine Andrade Aguiar, Flávia Cristina
Morone Pinto et al. "Curativo com filme e gel de
biopolímero de celulose bacteriana no
tratamento de feridas isquêmicas após
revascularização de membros inferiores.",
Revista do Colégio Brasileiro de Cirurgiões,
2019
Publication
-
- 28 clinicaltrials.gov <% 1
Internet Source
-

29

Saulo Nani Leite, Thiago Antônio Moretti de Andrade, Daniela dos Santos Masson-Meyers, Marcel Nani Leite et al. "Phototherapy promotes healing of cutaneous wounds in undernourished rats", *Anais Brasileiros de Dermatologia*, 2014

Publication

<% 1

30

Djamel Tahtat, Mohamed Mahlous, Samah Benamer, Assia Nacer Khodja, Souad Larbi Youcef, Nadjet Hadjarab, Wassila Mezaache. "Influence of some factors affecting antibacterial activity of PVA/Chitosan based hydrogels synthesized by gamma irradiation", *Journal of Materials Science: Materials in Medicine*, 2011

Publication

<% 1

31

Submitted to Universiti Teknologi Malaysia

Student Paper

<% 1

32

Pascual, Ana, Jeremy PK Tan, Alexander Y. Yuen, Julian M. W. Chan, Daniel J. Coady, David Mecerreyes, James L Hedrick, Yi Yan Yang, and Haritz Sardón. "Broad-Spectrum Antimicrobial Polycarbonate Hydrogels with Fast Degradability", *Biomacromolecules*, 2015.

Publication

<% 1

33

Submitted to University of Northumbria at Newcastle

Student Paper

<% 1

Vincenzo Guarino, Rosaria Altobelli, Francesca

- 34 della Sala, Assunta Borzacchiello, Luigi Ambrosio. "Chapter 4 Alginate Processing Routes to Fabricate Bioinspired Platforms for Tissue Engineering and Drug Delivery", Springer Science and Business Media LLC, 2018 <% 1
- Publication
-
- 35 jcdr.net <% 1
- Internet Source
-
- 36 www.o-wm.com <% 1
- Internet Source
-
- 37 Nandana Bhardwaj, Dimple Chouhan, Biman B. Mandal. "Tissue Engineered Skin and Wound Healing: Current Strategies and Future Directions", Current Pharmaceutical Design, 2017 <% 1
- Publication
-
- 38 Submitted to Loughborough University <% 1
- Student Paper
-
- 39 Submitted to University of Wales Swansea <% 1
- Student Paper
-
- 40 Submitted to CSU, Long Beach <% 1
- Student Paper
-
- 41 Submitted to A.T. Still University - Arizona <% 1
- Student Paper
-

- 42 Cynthia A. Fleck. "Wound Assessment Parameters and Dressing Selection", Advances in Skin & Wound Care, 09/2006 <% 1
Publication
-
- 43 Submitted to University of Greenwich <% 1
Student Paper
-
- 44 www.portalnepas.org.br <% 1
Internet Source
-
- 45 hdl.handle.net <% 1
Internet Source
-
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Internet Source
-
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Internet Source
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Internet Source
-
- 49 lib.jinr.ru <% 1
Internet Source
-
- 50 www.intechopen.com <% 1
Internet Source
-
- 51 Valencia, I.C.. "Chronic venous insufficiency and venous leg ulceration", Journal of the American Academy of Dermatology, 200103 <% 1
Publication
-

52

T Velnar, T Bailey, V Smrkolj. "The Wound Healing Process: An Overview of the Cellular and Molecular Mechanisms", Journal of International Medical Research, 2009

<% 1

Publication

53

Janice M. Burke. "Stimulation of DNA synthesis in human and bovine RPE by peptide growth factors: The response to TNF- α and EGF is dependent upon culture density", Current Eye Research, 2009

<% 1

Publication

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