CLINICAL MANAGEMENT

extra

Optimizing the Moisture Management Tightrope with Wound Bed Preparation 2015[©]





AMA PRA

2.5 Contact Hours

R. Gary Sibbald, BSc, MD, DSc (Hons), MEd, FRCPC (Med)(Derm), FAAD, MAPWCA • Professor • Medicine and Public Health • University of Toronto • Toronto, Ontario, Canada • Director • International Interprofessional Wound Care Course & Masters of Science in Community Health (Prevention & Wound Care) • Dalla Lana Faculty of Public Health • University of Toronto • Past President • World Union of Wound Healing Societies • Clinical Editor • Advances in Skin & Wound Care • Philadelphia, Pennsylvania

James A. Elliott, MS • Government Relations Director • Canadian Association of Wound Care • Knowledge Translation Research Director • Toronto Regional Wound Clinics • Toronto, Ontario, Canada

Elizabeth A. Ayello, PhD, RN, ACNS-BC, CWON, ETN, MAPWCA, FAAN • Faculty • Excelsior College of Nursing • Albany, New York • Senior Advisor • The John A. Hartford Institute for Geriatric Nursing • New York, New York • President • Ayello, Harris & Associates • New York, New York • Course Coordinator • International Interprofessional Wound Care Course • Clinical Editor • Advances in Skin & Wound Care • Philadelphia, Pennsylvania

Ranjani Somayaji, MD, BScPT, FRCPC • Clinical Lecturer • Division of Infectious Disease, Department of Medicine • Cumming School of Medicine • University of Calgary • Calgary, Alberta, Canada

Dr Sibbald has disclosed that he is a board member of Hollister; was a board of Coloplast and Systagenix/Acelity; is a consultant for Hollister; was a consultant for Coloplast and Systagenix/Acelity; is a recipient of grants from Systagenix/Acelity, Hollister, Healthpoint/Smith & Nephew, and Ferris Manufacturing; is a member of the speaker's bureau for Coloplast, Systagenix/Acelity, Hollister, and Molnlycke; receives payment for the development of educational programs from Coloplast, Systagenix/Acelity, and Hollister; and his spouse/partner (if any) has disclosed that he/she has no financial relationships with, or financial interests in, any commercial companies pertaining to this educational activity. Dr Ayello, Mr Elliott, and Dr Somayaji have disclosed that they have no financial relationships related to this article; and their spouses/partners (if any) have disclosed that they have no financial relationships with, or financial interests in, any commercial companies pertaining to this educational activity. Copyright 2015. R. Gary Sibbald.

All staff and planners, including spouses/partners (if any), in any position to control the content of this CME activity have disclosed that they have no financial relationships with, or financial interests in, any commercial companies pertaining to this educational activity.

Lippincott CME Institute has identified and resolved all conflicts of interest concerning this educational activity.

To earn CME credit, you must read the CME article and complete the quiz and evaluation on the enclosed answer form, answering at least 13 of the 18 questions correctly.

This continuing educational activity will expire for physicians on October 31, 2016, and for nurses on October 31, 2017.

If you need CME or CE STAT, take the test online at: http://cme.lww.com for physicians and www.nursingcenter.com for nurses. Complete CE/CME information is on the last page of this article.

PURPOSE

To provide an overview of moisture management and its importance in wound care.

TARGET AUDIENCE:

This continuing education activity is intended for physicians and nurses with an interest in skin and wound care.

OBJECTIVES:

After participating in this educational activity, the participant should be better able to:

- 1. Summarize causes and treatments for moisture balance issues of chronic wounds.
- 2. Recognize the properties of dressings used for treatment for moisture management of chronic wounds and antiseptic agent cytotoxicity.
- 3. Explain study findings of the effectiveness of dressing choices for treatment of chronic wounds.

ABSTRACT

OBJECTIVE: To provide an overview of moisture management and its importance in wound care. The authors evaluate the impact of moisture management for optimal wound care and assess current wound management strategies relating to antisepsis and moist wound healing utilizing the wound bed preparation paradigm 2015 update. The discussion distinguishes the form and function of wound care dressing classes available for optimal moisture management.

CONCLUSION: Moisture management for chronic wounds is best achieved with modern moist interactive dressings if the wound has the ability to heal.

KEYWORDS: moisture management, wound healing, antisepsis

ADV SKIN WOUND CARE 2015;28:466-76; quiz 477-8.

INTRODUCTION

A wound (ulcer) is a loss of epidermis with a dermal or deeper base, representing a disruption of skin integrity with tissue damage. Wounds can have vascular, traumatic, inflammatory, infectious, or malignant etiologies. Acute wound healing occurs along a concerted biochemical cascade. A wound can become chronic if the inflammatory or proliferative phases of the cascade stall. ^{1,2}

Distinct biochemical differences exist between healing and stalled chronic wounds. In healing wounds, cellular mitosis increases, whereas proinflammatory cytokines and matrix metalloproteinases decrease. In chronic wounds, the reverse process occurs. Following the same pattern, growth factors increase, and cellular response is rapid in healing wounds, whereas growth factor levels are suboptimal and cellular response senescent in chronic wounds.

Chronic wounds are prevalent and cause substantial morbidity, mortality, and increased healthcare costs.³ The wound bed preparation (WBP) paradigm provides a framework for care of chronic wounds, with an emphasis on an interprofessional approach. This article explores the use of WBP in chronic wound care. Moisture management will be discussed, including cleansing, antisepsis, and moist wound healing principles.

MOISTURE MANAGEMENT AND WOUND HEALING

Moisture management and moist wound healing concepts were established by the work of Winter⁴ in animal models and Hinman and Maibach⁵ in human models. Moist wound environments enhance wound healing and promote new tissue growth. In contrast, excess or insufficient moisture impairs the healing process and causes breakdown of the wound bed and surrounding skin. These tissue alterations increase the risk of bacterial damage from superficial critical colonization and deep/ surrounding wound infection.⁶ Low moisture levels may also lead to necrosis and eschar formation, hindering wound reepithelialization and closure. Thus, moisture balance of the wound bed is critical for wound healing.⁷

THE WOUND BED PREPARATION PARADIGM

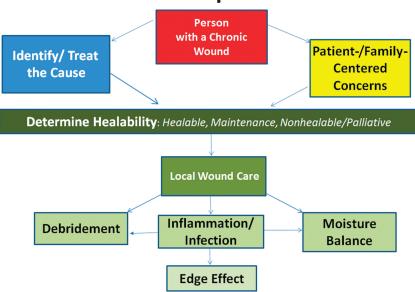
Wound Bed Preparation 2015 is a structured approach to wound healing. Building on previous editions, this WBP paradigm adds healability determination into the comprehensive assessment (Figure 1). This assessment should also identify patient-/family-centered concerns and an accurate diagnosis of wound etiology (ie, the wound cause [see Table 1]). The 3 components of local wound care—debridement, inflammation/infection, and moisture balance management—should be addressed after completing the comprehensive patient assessment, including the division of wounds into healable, maintenance, and nonhealable healing potential categories. The clinician should distinguish:

- **healable wounds** with adequate blood supply that can be healed if the underlying cause is addressed.
- *maintenance wounds* have healing potential, but also have patient or health system barriers compromising healing, including patient nonadherence to treatment or healthcare resource limitations.
- *nonhealable wounds* (including palliative wounds) cannot heal because of irreversible causes or associated illnesses, including critical ischemia or nontreatable malignancy.

In maintenance and nonhealable wounds, a relatively conservative approach should be taken, potentially involving conservative debridement of slough, bacterial reduction through antisepsis, and moisture reduction (Table 2).

Figure 1.
WOUND BED PREPARATION PARADIGM 2015

Wound Bed Preparation 2015



In healable wounds (Table 2), there are 3 initial local wound care components that should be addressed:

- *debridement* of necrotic tissue that may include active surgical removal of debris to bleeding tissue;
- *inflammation/infection* recognition and management, followed by topical and systemic therapies as appropriate; and
- moisture balance in the wound bed interface.

See Table 2 for a summary of local wound care strategies.

HEALABLE WOUNDS: AN APPROACH TO MOISTURE-BALANCE DRESSINGS

Moisture balance at the wound bed interface may be achieved with a variety of dressings (Table 3). There are 5 major choices of antimicrobial dressings (silver, polyhexamethylenebiguanide [PHMB], iodine, methylene blue/crystal violet, and honey), with 2 of these choices having anti-inflammatory properties (silver, honey). Moisture-balance dressing classes are often combined with antibacterial and anti-inflammatory dressings in healable wounds to manage inflammation/infection according to the clinical characteristics of the wound. Validated tools that may be utilized to diagnose wound infection/inflammation before using these dressings include the "NERDS" and "STONEES" criteria and SIBBALD cubes^{6,11} (Figure 2).

Table 1. TREATMENT OF THE CAUSE FOR SELECTED COMMON CHRONIC WOUNDS

Wound Type	Treatment of the Cause
Venous ulcers	Bandages for healing
	Stockings to prevent recurrence
Pressure ulcers	Redistribute pressure (relieve heel pressure)
	• Promote physical activity as tolerated
	Manage incontinence and moisture
	Reduce shear
	Enhance and optimize nutrition
Diabetic (neurotrophic) foot ulcers	Vascular: ensure adequate vascular supply
Callus = pressure	Infection: control superficial critical
Blister = friction	colonization/deep + surrounding infection
and shear	 Redistribute plantar pressure

Table 2. SUMMARY OF LOCAL WOUND CARE STRATEGIES BY WOUND HEALABILITY CLASSIFICATION

Wound Healability Classification	Surgical Debridement	Inflammation/Infection Management	Moisture Management
Healable	Active	Treat inflammation/infection (topically or systemic) + antisepsis as required	Moisture balance
Maintenance	Conservative (no bleeding)	Bacterial reduction—antisepsis	Moisture reduction
Nonhealable	Comfort removal of slough	Bacterial reduction—antisepsis	Moisture reduction

Using the NERDS mnemonic, if 3 or more are present, treat topically:

Nonhealing wounds Exudative wounds

Red and bleeding wound surface granulation tissue

Debris (yellow or black necrotic tissue) on the wound surface

 ${f S}$ mell or unpleasant odor from the wound

Using the STONEES mnemonic, if 3 or more are present, treat systemically:

Size is bigger

Temperature of 3° F or more versus mirror image

Os (probe to or exposed bone) New or satellite areas of breakdown

Exudate is increased

Table 3. SUMMARY OF MODERN DRESSING CATEGORIES ORDERED BY INCREASING ABSORBENCY

Modern Dressing Category	Comment	Average Wear Time
Hydrogels ^a	Contain 70%-90% moisture	1–3 d
Donate moisture	Donates moisture to the wound	
	Bioresorbable	
	Can be combined with silver, iodine (cadexomer) for antimicrobial action	
Films ^a	Protective layer	3–7 d
Moisture neutral	Does not donate or absorb a large amount of exudate	
Hydrocolloids ^a	Water-binding and water-repelling components	2-7 d
	Will absorb small to moderate amount of moisture	
Hydrofibers	Bind small to moderate amount of exudate	1–3 d
	Fluid lock, nonbioresorbable	
	Can be combined with silver for antimicrobial action	
Calcium alginates ^a	Absorb small to moderate amounts of exudate onto outer surface of dressing	1–3 d
	Fibers are bioresorbable, releasing calcium (hemostasis property) and resorbing sodium to form a hydrogel with exudate fluid	
	Can be combined with silver and honey for antibacterial action	
Foams	Absorb moderate amounts of exudate	2-7 d
	Fluid balance with the dressing giving back some exudate that prevents wound surface from dehydrating	
	Can be a method of delivering an antibacterial agent (silver) or containing a nonrelease antibacterial agent for antibacterial action above the wound surface (PHMB, methylene blue/gentian violet)	
Superabsorbents	Absorb a larger amount of exudate	1–3 d
	Fluid lock technology equivalent to diapers	

Figure 2.
SIBBALD CUBE

© 2013 IIWCC Non-healing Superficial: •Exudate↑ **NERDS** criteria •Red + Bleeding •Debris •Smell Size is bigger •Temperature ↑ Deep: Os (probes, exposed) STONEES criteria •New breakdown Any 3 need Physician to •Exudate↑ ler systemic •Erythema , Edema Based on clinical criteria

Sibbald Cube[©]

- Use in the Healable Wound that is "stalled". (Less than 30% decrease in size in 4 weeks)
- Correct & modify cofactors using Wound Bed Preparation[®]

*Test to detect high surface MMPs or based on clinical criteria

SUPERFICIAL	HIGH PROTEASES *	LOW PROTEASES
HIGH BACTERIA ≥ 3 NERDS	Anti-inflammatory +Antimicrobial dressing	Antimicrobial dressing
LOW BACTERIA < 3 NERDS	Anti-inflammatory dressing	Moisture-balance dressing

DEEP & SURROUNDING TISSUE	HIGH PROTEASES *	LOW PROTEASES
HIGH BACTERIA ≥ 3 STONEES	Oral anti-inflammatory antimicrobial	Systemic antimicrobial
≥ 5 STUNEES		
LOW BACTERIA	Systemic anti-	No systemic therapy
< 3 STONEES	inflammatory	

Erythema and/or edema (cellulitis) Smell

MOISTURE-BALANCE DRESSINGS

The Cochrane reviews (Table 4) state there is often no current evidence to support the effectiveness of many of these dressings over a comparator dressing or standard wound care. Yet, modern moist interactive dressings can offer several advantages. Gauze dressings, for example, need frequent changes (1–3 times per day). This can result in intensive demand for nursing care. Furthermore, gauze dressings are associated with increased patient pain and potential for wound trauma upon removal. ¹² Although gauze is relatively inexpensive, the costs of nursing services and patient time required for frequent changes can sometimes make gauze less cost-effective than most modern dressings, particularly when these dressings are used appropriately.

The following sections (Table 3) will discuss moisture-balance dressing categories. Particular attention will be given to chemical composition, form, function, and clinical application.

Hydrogels

6a

Hydrogel (ie, hydrated polymer) dressings have a high water content (60%–90%). Hydrogels are capable of providing moisture. This feature aids the autolytic debridement of sloughy or necrotic wound tissue. Hydrogels are clear or translucent, vary in viscosity, and are available in 3 forms: amorphous (most common), impregnated gauze, and as a wafer. In clinical settings, the high water content can lead to periwound maceration. To prevent this moisture-associated damage, a periwound barrier should be applied after wound cleansing (saline or water preferred). Four barriers are available: using film-forming liquid acrylate spray or wipe, zinc oxide ointment, petrolatum, or a windowed occlusive dressing (film or hydrocolloid).

Films

Film dressings are transparent polyurethane dressings with or without adhesives. They are often used for local protection of a wound at the late re-epithelialization stage or to protect a recently healed wound. The choice of a nonadherent versus a film dressing

Table 4.
SUMMARY OF RECENT LITERATURE ON MOIST BALANCE DRESSINGS

Dressing(s)	Brief Summary ^a
Alginates: diabetic neuropathic neuroischemic/ischemic foot ulcers	A 2013 Cochrane review investigated alginate dressings for healing diabetic neuropathic/ischemic/neuroischemic foot ulcers. ²⁵ It included 6 studies (n = 375) that compared alginate dressings with basic wound contact dressings, foam dressings, and silver-containing, fibrous-hydrocolloid dressings.
	Conclusion: No evidence found to suggest alginate wound dressings are more effective at healing diabetic foot ulcers than other dressings. Small sample size merits caution.
Alginates: in pressure ulcers	A 2015 Cochrane review investigates alginate dressings for healing pressure ulcers. ²⁶ It included 6 studies (n = 336) that compared alginate dressings with hydrocolloid dressings, silver-containing alginate dressings, and radiant heat therapy.
	Conclusion: The relative effects of alginate dressings compared with alternative treatments are unclear.
Alginates: chronic ulcers in older adults (>65 y)	A 2015 Canadian Health Technology Assessment (HTA) investigated prevention and treatment of chronic ulcers in the older adults. ²¹
	Conclusion: Dressings containing calcium alginate may lead to shorter healing time of pressure ulcers in older adults.
Alginates and films: split-thickness skin grafts	It also concluded healing effects of other dressings in this specific age group are insufficiently studied. A 2013 randomized controlled trial (RCT) ($n = 38$) by Läuchli et al ²⁴ compared calcium alginate dressings vs polyurethane film dressing for split-thickness skin graft donor sites.
	Conclusion: Film dressings resulted in initial lower pain scores, whereas alginate dressings were found to cause fewer additional dressing changes and less leakage.
Foams: diabetic neuropathic neuroischemic/ischemic foot ulcers	A 2013 Cochrane review investigated foam dressings for healing diabetic foot ulcers ²⁷ ; included 6 studies (n = 157) that compared foam dressings with basic wound dressings.
	Conclusion: No evidence found to suggest foam wound dressings are more effective in healing diabetic foot ulcers than other dressings. Small sample size merits caution.
Foams: venous leg ulcers	A 2013 Cochrane review investigated foam dressings for venous leg ulcers. 28 It included 12 studies (n = 1023) that evaluated the effects of any type of foam dressing in the treatment of venous ulcers.
	Conclusion: The current evidence base does not suggest that foam dressings are more effective in the healing of venous leg ulcers than other wound dressing treatments.
Hydrocolloids: diabetic neuropathic neuroischemic/ischemic foot ulcers	A 2013 Cochrane review investigated hydrocolloids for healing diabetic foot ulcers. ²⁹ It included 5 studies (n = 535) that compared hydrocolloid dressings with basic wound contact dressings, foams, silver, and a topical cream containing plant extracts.
	Conclusion: No evidence found to suggest that any type of hydrocolloid wound dressing is more effective in healing diabetic foot ulcers than other types of dressing or a topical cream containing plant extracts. Small sample size merits caution.
Hydrocolloids: nonadherent vs traditional dressings for wounds in general	A 2011 Canadian HTA investigated hydrocolloid vs traditional dressings. ²² It reviewed 1 relevant HTA report, 3 relevant systematic reviews, and 7 relevant evidence-based guidelines.
	Conclusion:
	 Evidence suggests that hydrocolloid dressings may be clinically effective for venous leg and pressure ulcers.
	 Several guidelines recommend nonadherent dressings for venous leg and pressure ulcers, amputations, and chronic wounds.
Hydrofibers and alginates for nonischemic diabetic foot ulcers	A 2007 prospective RCT (n = 67) by Jude et al ²⁰ compared hydrofiber dressing containing ionic silver vs calcium alginate dressings in nonischemic diabetic foot ulcers.
	Conclusion: When added to standard care with appropriate off-loading, hydrofiber dressings containing ionic silver were associated with favorable clinical outcomes compared with calcium alginates, specifically in ulcer depth reduction and in infected ulcers requiring antibiotic treatment. (Note that the hydrofiber had silver, and the comparator calcium alginate did not.) (continues)

Table 4.
SUMMARY OF RECENT LITERATURE ON MOIST BALANCE DRESSINGS, CONTINUED

Brief Summary ^a
A 2015 Cochrane review investigated hydrogel dressings for healing pressure ulcers. ³⁰ It included 11 studies (n = 523) that compared hydrogel dressings with alternative dressings or no dressings for pressure ulcers Stage II or higher.
Conclusion: It is not clear if hydrogel dressings are more or less effective than other treatments in healing pressure ulcers or if different hydrogels have different effects.
A 2013 Cochrane review investigated hydrogel dressings for healing diabetic foot ulcers. ¹⁸ It included 5 studies (n = 446) that compared hydrogel dressings with alternative dressings or no dressings.
Conclusion: There is some evidence to suggest that hydrogel dressings are more effective in healing (lower grade) diabetic foot ulcers than basic wound contact dressings.
There is no evidence to suggest that hydrogels are more effective than larval therapy or platelet-derived growth factors in healing diabetic foot ulcers, nor that 1 brand of hydrogel is more effective than another in ulcer healing.
A 2010 Cochrane review investigated hydrogels for debridement of diabetic foot ulcers. ¹⁹ It included 6 RCTs comparing hydrogel with gauze or standard care.
Conclusion: There is evidence to suggest that hydrogels increase the healing rate of diabetic foot ulcers compared with gauze dressings or standard care.
A 2014 Canadian HTA investigated debridement procedures for diabetic foot ulcers. ²³ It reviewed 12 relevant studies and 7 guidelines.
Conclusion: Autolytic (hydrogel) debridement and enzymatic debridement (clostridial collagenase ointment) are more clinically effective for wound debridement procedures for the treatment of diabetic foot ulcers than standard wound care.

with adhesive backing is partly determined by the fragility of the surrounding skin. Film dressings with acrylic adhesives can cause skin tears in contrast to silicone-coated films that decrease pain and trauma with dressing removal. Film materials are semi-occlusive, have relatively no absorptive capacity, and have a varying degree of permeability (referred to as the moisture vapor transmission rate) that allow for differential evaporation of the water molecules through the dressing. Remove acrylic adhesives by gently pulling laterally in a repeated clockwise rotation.

Hydrocolloids

Hydrocolloids are most commonly available in a wafer type of occlusive dressing that consist of gel-forming agents (containing carboxymethylcellulose) with a flexible, water-resistant outer layer. The dressings have an adhesive and come in a variety of shapes designed for body areas, including the sacrum and heels. Hydrocolloid dressings are mildly absorptive and have a wear time equivalent to foam dressings (up to 7 days) but longer than most other dressing classes. For application of hydrocolloid dressings, the wound margin should be overlapped by 1 to 2 cm to form an

adhesive seal. This overlap also prevents exudate leakage from the edges of the dressing. When these dressings are used for autolytic debridement, they may need to be changed more frequently. Removal of nonviable slough from the surface of the wound may also be required at dressing change to prevent odor or secondary bacterial proliferation under the dressing.

Hydrofibers

Hydrofiber dressings consist of carboxymethylcellulose spun into a fiber format instead of the gelled form in hydrocolloid dressings. The fiber gives the dressing tensile strength, and it can usually be removed easily in 1 piece. As the spun hydrofibers bind exudate with interior fluid lock, the dressing promotes very little autolytic debriding. As the dressing absorbs fluid, the hydrofibers are converted into a gel. Hydrofiber dressings are thin and have low to moderate absorbency, although thicker newer dressing options have increased absorbency. These dressings need a secondary dressing to keep them in place because the addition of an adhesive will interfere with the fluid absorption properties of the dressing.

Calcium Alginates

Calcium alginates are nonwoven biodegradable fibers processed from acids derived from brown seaweed. When calcium alginates bind fluid as in a wound, the calcium ions are donated to the wound surface, and the absorbed sodium results in the formation of a soluble hydrogel. Calcium alginate dressings are able to absorb up to 20 times their weight in fluid. Once in gel form, the dressings can promote autolytic debridement of the wound. Uniquely to calcium alginate dressings, release of calcium ions into the wound bed can also help in hemostasis without the formation of hemorrhagic crust on the wound surface. These dressings are manufactured in sheets (lateral fluid wicking) or in ropes (vertical fluid wicking) and can readily conform to wounds of varying shapes. Alginate dressings are bioresorbable and need a secondary dressing with similar application principles for hydrogels or hydrofibers. If any alginate fibers are left intact at dressing change, they can be moistened to dissolve and do not have to be removed mechanically. If the fibers remain dry, a water-donating hydrogel may be a better dressing choice.

Foams

Foam dressings are manufactured most commonly as polyurethane foams. These dressings absorb a moderate to large amount of exudate. Foam dressings can consist of 2 to 3 layers with a hydrophilic contact surface between the foam and a hydrophobic backing. Foam dressings are manufactured with normal absorbency, light or less absorptive dressings, with and without adhesives or borders. The silicone adhesive format has demonstrated decreased pain on dressing removal compared with the more traditional acrylic adhesives. 12 Exudate absorbs into hollow polyurethane pores, creates equilibrium, and donates moisture back to the wound with increasing saturation to achieve a fluid balance. The fluid exchange function can lead to periwound maceration. Some of the more advanced foams have variable pore sizes that lead to partial fluid retention in addition to the traditional fluid exchange functions. Periwound maceration can also be minimized if a periwound barrier is applied, and the foam is cut to the wound size, fenestrated on the top to wick to a secondary superabsorbent dressing, or changed more frequently. An alternate foam core with polyvinyl alcohol can provide autolytic debridement not provided by the traditional polyurethane core. Foams have also been combined with antiseptics (eg, silver, PHMB, methylene blue/crystal violet) and other agents to serve as a delivery vehicle for active therapies at the wound surface.

Superabsorbents

Superabsorbent polymer–containing wound dressings are best suited to manage highly exudative wounds. ¹³ These dressings can absorb an enormous amount of water relative to their dry weights.

Superabsorbent polymers are the same technology utilized in diapers, feminine hygiene materials, and adult incontinence products. ¹⁴ Superabsorbent dressings are typically manufactured from acrylic acid. They undergo polymerization by suspension or crosslinking, which accounts for their absorptive and protein-binding properties (ie, proteases). ^{15,16} They have multiple layers, a large absorbent surface, a fluid lock to prevent periwound maceration, and a contact layer that protects the wound base from the inner core that can become saturated with wound exudate. The core fluid locking materials may include powders, crystals, or gelling agents that work by osmosis, with fibers having a capillary-like action. Secondary dressings are necessary for fixation to the surface of the wound if there is no adhesive.

Recent Literature on Moisture Balance Dressings

The authors searched The Cochrane Library, Ovid MEDLINE, University of York Centre for Reviews and Dissemination database, and Google Scholar for systematic reviews, health technology assessments, and high-quality randomized controlled trials published from January 2007 to June 2015. The terms "foam," "superabsorbent," "calcium alginate," "hydrogel," "acrylate," "hydrocolloid," or "film" and word variations of these were searched. The authors used the GRADE system to assess the quality of articles. ¹⁷ Hand referencing was also utilized. The results of the literature review are displayed in Table 4. Most dressing versus dressing comparisons did not yield definitive conclusions. However, some exceptions exist, including the following:

- \bullet hydrogels over basic contact wound dressings for diabetic foot ulcers 18
- \bullet hydrogels over gauze or standard care for debridement of diabetic foot ${\rm ulcers}^{19}$
- hydrofiber dressing containing ionic silver over calcium alginate dressings in nonischemic diabetic foot ulcers²⁰ (the comparator was not equal because it did not contain silver)
- calcium alginate dressing over other comparator treatments for pressure ulcers (PrUs) in older adults²¹
- \bullet hydrocolloid dressings over other comparator treatments for venous leg and pressure ulcers 22
- autolytic (hydrogel) and enzymatic debridement (clostridial collagenase ointment) for debridement of diabetic foot ulcers over standard wound care²³
- \bullet calcium alginate over polyure thane film dressing for split-thickness skin graft donor sites. 24

Wound Cleansing

The goal of wound cleansing is to promote healing through improved wound assessment, increased comfort with adherent dressing removal, and possible rehydration of the wound bed. Antiseptics have been used as wound cleansing agents for decades,

but questioned in recent years because of a paucity of evidence for their use. The standard of care for wound cleansing is to use solutions that are as gentle and noncytotoxic to the wound as possible, such as saline, water, or acetic acid (0.5%–1.0%). A compress is when these solutions are applied to gauze, and the excess is rung out before application. The result is a net movement of fluid from the wound surface to the gauze via astringent (coagulate protein) action. A soak uses the same procedure of saturating gauze, but the gauze is then applied saturated, resulting in a net fluid movement into a dry wound surface from the gauze.

An updated Cochrane Collaboration review in 2013 for cleansing PrUs concluded, "There is no good trial evidence to support use of any particular wound cleansing solution or technique for pressure ulcers." These same principles apply to irrigation. This technique can cause more harm than benefit if the force applied causes more pain or tissue damage. If the bottom of the wound is not visualized, and irrigation fluid remains behind, it may also form the nidus for bacterial abscess.

MAINTENANCE AND NONHEALING WOUNDS

A conservative approach should be taken for the management of wounds with compromised healing potential. The focus should be placed on patient-concerned concerns, especially pain and optimizing activities of daily living. Antiseptics are frequently used for the purposes of moisture reduction and control of bacterial burden. As with healable wounds, solutions with minimal potential for cytotoxicity should be utilized. Some antiseptic solutions are more cytotoxic to fibroblasts than other solutions, and although toxicity is often less in vivo, the impact may be increased in nonhealing wounds.

Most antiseptics are bactericidal with a broad spectrum of action. They often have many targets, including cell walls, cell membranes, cytoplasmic organelles, and DNA. ³² Bacterial resistance of antiseptics is very low, and their use is preferred topically compared with topical antibiotics. Topical antibiotic use should be generally avoided to lower the risk of bacterial resistance and adverse effects. For maintenance and nonhealing wounds, systemic antibiotics are reserved for deep and surrounding infections.

In an area of inadequate blood supply or uncontrolled edema (eg, congestive heart failure, refractory venous disease), moisture reduction and the use of topical antisepsis with or without a secondary dressing may be beneficial. For example, in persons with distal gangrene, antiseptic agents with low toxicity may be used, including 10% povidone-iodine, chlorhexidine, or its derivative PHMB. The agents are best applied around the proximal edges of the gangrene to decrease the risk of infection and prevent tissue breakdown at the edge between the gangrenous and viable tissue.

Active or aggressive debridement that creates bleeding is not recommended in maintenance and nonhealing wounds. The rea-

son is that aggressive debridement further compromises tissue, leading to potential deep infection. For example, in diabetic neurotropic foot ulcers with inadequate vascular supply, active debridement leads to bleeding, further callus formation, and an expanding ulcer. Conservative debridement with callus removal followed by local wound care as previously discussed is recommended. Similarly, in PrUs without healing potential, the same wound care principles are recommended, along with application of strategies to minimize pressure and shear forces.

Table 5 lists the common antiseptics in ascending order of tissue toxicity. Although the "red" agents have increased potential for cytotoxicity, they may be useful in specific circumstances.

Chlorhexidine, PHMB, and povidone-iodine have their antibacterial activity by attacking bacterial cell walls, cytoplasmic organelles, or nucleic acids. Dilute acetic acid (0.5%–1%) lowers the surface pH of wounds. This has antipseudomonal activity as *Pseudomonas* species grow best in alkaline pH environments. White vinegar (5% acetic acid) can be diluted 1:5 to 1:10 with potable or sterile water and applied locally as an alternating compress. Gauze can be moistened with the acetic acid and squeezed to remove excess moisture. The gauze is then placed on the wound for 30 to 60 seconds and discarded. A second gauze application follows for 5 to 10 minutes. Although *Pseudomonas aeruginosa* can often be found in chronic wounds, guidelines have increasingly suggested it seldom requires systemic treatment in the absence of deep and surrounding *Pseudomonas* predominant infection.³³

Antiseptics with high potential for cytotoxicity include dyes, bleaches, hydrogen peroxide, and quaternary ammonium compounds. Dyes including agents such as scarlet red and mercurochrome

Table 5.
SELECT ANTISEPTIC AGENTS LISTED BY
INCREASING CYTOTOXICITY

Agent	Effects
Chlorhexidine or PHMB	Low toxicity
Povidone-iodine (Betadine)	Broad spectrum
Acetic acid—vinegar diluted 1:5 to 1:10	Pseudomonas
Saline/sterile water	Not antibacterial
Dyes—scarlet red, proflavine	Select out gram negative
Sodium hypochlorite—Dakin solution, EUSOL	Toxic = bleach
Hydrogen peroxide	Action = fizz
	Very high toxicity

Agents are color coded by safety profile and antiseptic action: green = low toxicity potential, yellow = no antibacterial effect, red = high toxicity potential.

are more active against gram-positive than gram-negative bacteria. Bleach (sodium hypochlorite) is an excellent external environmental agent, often used to decrease bacterial contamination on working surfaces and objects. Bleach is also prepared as Dakin solution or Edinburgh University solution of lime (EUSOL).

Patients with extensive wounds and adherent, difficult-to-remove dressings can be clinically challenging. Soaking each individual wound for 5 to 10 minutes or removing the dressings in the bathtub may help reduce dressing removal pain and trauma. A dilute acetic solution will acidify water and help decrease bacteria. Bleach (5–10 mL in 5 L) with acidification releases hypochlorous acid and also acts as an astringent to coagulate protein. These processes may decrease the bacterial burden and be beneficial if used sparingly. 34

Hydrogen peroxide has a broad range of activity, with a short period of antibacterial action on the skin. It is active only when fizzing and is associated with air emboli if used in deep cavities. ³⁵ Lastly, quaternary ammonium compounds have detergent-like actions with a broad antimicrobial activity but have a higher level of tissue toxicity than other agents. Therefore, quaternary ammonium compounds are not a recommended agent in wound management.

SUMMARY

Moisture management for chronic wounds is best achieved with modern moist interactive dressings if the wound has the ability to heal. For nonhealable or maintenance wounds, moisture reduction, bacterial reduction, and conservative debridement of slough are recommended. Each patient must be considered individually, and wounds assessed for pain, local wound fragility, and tissue viability in order to make the best choice for local wound care utilizing the WBP paradigm.

PRACTICE PEARLS

- All chronic wounds should be classified as healable, non-healable, or maintenance.
- Moisture-balance dressings are important for healable wounds, with moisture reduction often more appropriate for nonhealable or maintenance wounds.
- Sharp surgical debridement is appropriate for healable wounds, with conservative surgical debridement of slough more important for the nonhealable and maintenance wound.
- Critically colonized wounds (≥3 NERDS criteria) require antimicrobial dressings, with deep and surrounding infections (≥3 STONEES criteria) most appropriately treated with systemic antimicrobial agents.
- There is very little scientific evidence for wound cleansing, and each patient should be evaluated to ensure the technique results in more benefits than the amount of associated pain, or tissue damage, including retained fluid in deep cavities.

REFERENCES

- 1. Singer AJ, Clark RA. Cutaneous wound healing. N Engl J Med 1999;341:738-46.
- Schultz GS, Mast BA. Molecular analysis of the environments of healing and chronic wounds: cytokines, proteases and growth factors. Wounds 1998;10(6 Suppl F):1F-9F.
- Sen CK, Gordillo GM, Roy S, et al. Human skin wounds: a major and snowballing threat to public health and the economy. Wound Repair Regen 2009;17:763-71.
- Winter G. Formation of the scab and the rate of epithelization of superficial wounds in the skin of the young domestic pig. Nature 1962;193:293-4.
- Hinman CD, Maibach H. Effect of air exposure and occlusion on experimental human skin wounds. Nature 1963;200:37-8.
- Sibbald RG, Woo K, Ayello EA. Increased bacterial burden and infection: the story of NERDS and STONEES. Adv Skin Wound Care 2006;19:447-61.
- Okan D, Woo K, Ayello EA, Sibbald G. The role of moisture balance in wound healing. Adv Skin Wound Care 2007;20:39-53.
- Falanga V. Classifications for wound bed preparation and stimulation of chronic wounds. Wound Repair Regen 2000;8:347-52.
- Sibbald RG, Goodman L, Woo KY, et al. Special considerations in wound bed preparation 2011: an update. Adv Skin Wound Care 2011;24:415-36.
- Sibbald RG, Ovington LG, Ayello EA, Goodman L, Elliott JA. Wound bed preparation 2014 update: management of critical colonization with a gentian violet and methylene blue absorbent antibacterial dressing and elevated levels of matrix metalloproteases with an ovine collagen extracellular matrix dressing. Adv Skin Wound Care 2014; 27(3 Suppl 1):1-6.
- Woo K, Sibbald R. A cross-sectional validation study of using NERDS and STONEES to assess bacterial burden. Ostomy Wound Manage 2009;55(8):40-8.
- World Union of Wound Healing Societies. Principles of Best Practice: Minimising Pain at Wound Dressing-Related Procedures. A Consensus Document. London, UK: MEP Ltd; 2004:1-10
- Faucher N, Safar H, Baret M, Philippe A, Farid R. Superabsorbent dressings for copiously exuding wounds. Br J Nurs 2012;21(12):S22, S24, S26-S28.
- Pytlik E, Molino D, Moritz J. Superabsorbent polymers (SAP). In: Introduction to Polymers. Buffalo, NY: UB Engineering–University at Buffalo; 2005. www.eng.buffalo.edu/courses/ce435/Diapers/Diapers.html. Last accessed July 22, 2015.
- Wiegand C, Abel M, Ruth P, Hipler UC. Superabsorbent polymer-containing wound dressings have a beneficial effect on wound healing by reducing PMN elastase concentration and inhibiting microbial growth. J Mater Sci Mater Med 2011;22: 2583-90.
- Weigland C, Hipler C. A superabsorbent polymer-containing wound dressing efficiently sequesters MMPs and inhibits collagenase activity in vitro. J Mater Sci Mater Med 2013; 24:2473-8.
- Guyatt GH, Oxman AD, Vist GE, et al. GRADE: an emerging consensus on rating quality of evidence and strength of recommendations. BMJ 2008;336(7650):924-6.
- Dumville JC, O'Meara S, Deshpande S, Speak K. Hydrogel dressings for healing diabetic foot ulcers. Cochrane Database Syst Rev 2013;7:CD009101.
- Edwards J, Stapley S. Debridement of diabetic foot ulcers. Cochrane Database Syst Rev 2010;1:CD003556.
- Jude EB, Apelqvist J, Spraul M, Martini J, Silver Dressing Study Group. Prospective randomized controlled study of Hydrofiber dressing containing ionic silver or calcium alginate dressings in non-ischaemic diabetic foot ulcers. Diabet Med 2007;24:280-8.
- Swedish Council on Health Technology. Chronic Ulcers in the Elderly—Prevention and Treatment. Stockholm, Sweden: Swedish Agency for Health Technology Assessment and Assessment of Social Services; 2014.
- 22. Canadian Agency for Drugs and Technologies in Health. Non-adherent Versus Traditional Dressings for Wound Care: Comparative Effectiveness, Safety, and Guidelines. Ottawa, ON, Canada: Canadian Agency for Drugs and Technologies in Health; 2011:1-14.
- Canadian Agency for Drugs and Technologies in Health. Debridement Procedures for Managing Diabetic Foot Ulcers: A Review of Clinical Effectiveness, Cost-effectiveness, and Guidelines. Ottawa, ON, Canada: Canadian Agency for Drugs and Technologies in Health; 2014:1-38.
- Läuchli S, Hafner J, Ostheeren S, Mayer D, Barysch MJ, French LE. Management of split-thickness skin graft donor sites: a randomized controlled trial of calcium alginate versus polyurethane film dressing. Dermatology 2013;227:361-6.
- Dumville JC, O'Meara S, Deshpande S, Speak K. Alginate dressings for healing diabetic foot ulcers. Cochrane Database Syst Rev 2013;6:CD009110.

- Dumville JC, Keogh SJ, Liu Z, Stubbs N, Walker RM, Fortnam M. Alginate dressings for treating pressure ulcers. Cochrane Database Syst Rev 2015;5:CD011277.
- Dumville JC, Deshpande S, O'Meara S, Speak K. Foam dressings for healing diabetic foot ulcers. Cochrane Database Syst Rev 2013;6:CD009111.
- O'Meara S, Martyn-St James M. Foam dressings for venous leg ulcers. Cochrane Database Syst Rev 2013;5:CD009907.
- Dumville JC, Deshpande S, O'Meara S, Speak K. Hydrocolloid dressings for healing diabetic foot ulcers. Cochrane Database Syst Rev 2013;8:CD009099.
- Dumville JC, Stubbs N, Keogh SJ, Walker RM, Liu Z. Hydrogel dressings for treating pressure ulcers. Cochrane Database Syst Rev 2015;2:CD011226.
- Moore ZE, Cowman S. Wound cleansing for pressure ulcers. Cochrane Database Syst Rev 2013;3:CD004983.
- McDonnell G, Russell AD. Antiseptics and disinfectants: activity, action, and resistance.
 Clin Microbiol Rev 1999;12:147-79.
- Lipsky BA, Berendt AR, Cornia PB, et al. 2012 Infectious Diseases Society of America clinical practice guideline for the diagnosis and treatment of diabetic foot infections. Clin Infect Dis 2012;54(12):e132-73.
- 34. Mellerio J. Infection and colonization in epidermolysis bullosa. Dermatol Clin 2010;28:267-9.
- Brayfield A, ed. Martindale: The Complete Drug Reference. 38th ed. London, UK: Pharmaceutical Press: 2014.

Notice: Online CE Testing Only Coming in 2016!

Starting with the first issue of 2016, all CE tests must be completed online at CEconnection.com/aswc for nurses or at CME.LWW.com for physicians. Simply select the CE article you are interested in. Both the article and the test will available there. Mailing or faxing tests will no longer be options.

If you haven't done so already, you will want to create a user account for yourself – it's free to do so! Look for the Login link in the upper right hand corner of the screen.

For more than 120 additional continuing education articles related to skin and wound care topics, go to NursingCenter.com/CE.

GE COMECTION

CONTINUING MEDICAL EDUCATION INFORMATION FOR PHYSICIANS

Lippincott Continuing Medical Education Institute, Inc. is accredited by the Accreditation Council for Continuing Medical Education to provide continuing medical education for physicians.

Lippincott Continuing Medical Education Institute, Inc. designates this journal-based CME activity for a maximum of 1 AMA PRA Category 1 CreditTM. Physicians should only claim credit commensurate with the extent of their participation in the activity.

PROVIDER ACCREDITATION INFORMATION FOR NURSES

Lippincott Williams & Wilkins, publisher of the Advances in Skin & Wound Care journal, will award 2.5 contact hours for this continuing nursing education activity.

LWW is accredited as a provider of continuing nursing education by the American Nurses Credentialing Center's Commission on Accreditation.

This activity is also provider approved by the California Board of Registered Nursing, Provider Number CEP 11749 for 2.5 contact hours. LWW is also an approved provider by the District of Columbia, Georgia, and Florida CE Broker #50-1223. Your certificate is valid in all states.

OTHER HEALTH PROFESSIONALS

This activity provides ANCC credit for nurses and AMA PRA Category 1 CreditTM for MDs and DOs only. All other healthcare professionals participating in this activity will receive a certificate of participation that may be useful to your individual profession's CE requirements.

CONTINUING EDUCATION INSTRUCTIONS

- Read the article beginning on page 466.
- Take the test, recording your answers in the test answers section (Section B) of the CE enrollment form. Each question has only one correct answer.

- Complete registration information (Section A) and course evaluation (Section C).
- Mail completed test with registration fee to: Lippincott Williams & Wilkins, CE Group,
 74 Brick Blvd, Bldg 4 Suite 206, Brick, NJ 08723.
- Within 3 to 4 weeks after your CE enrollment form is received, you will be notified
 of your test results.
- If you pass, you will receive a certificate of earned contact hours and an answer key. Nurses who fail
 have the option of taking the test again at no additional cost. Only the first entry sent by
 physicians will be accepted for credit.
- A passing score for this test is 13 correct answers.
- NURSES: NEED CE STAT? Visit www.nursingcenter.com for immediate results, other CE activities
 and your personalized CE planner tool. No Internet access? Call 1-800-787-8985 for other
 rush service options.
- PHYSICIANS: NEED CME STAT? Visit http://cme.lww.com for immediate results, other CME activities, and your personalized CME planner tool.
- Questions? Contact Lippincott Williams & Wilkins: 1-800-787-8985.
 Registration Deadline: October 31, 2017 (nurses); October 31, 2016 (physicians).

PAYMENT AND DISCOUNTS

- The registration fee for this test is \$24.95 for nurses; \$22 for physicians.
- Nurses: If you take two or more tests in any nursing journal published by LWW and send in your CE enrollment forms together by mail, you may deduct \$0.95 from the price of each test. We offer special discounts for as few as six tests and institutional bulk discounts for multiple tests.
 Call 1-800-787-8985 for more information.