Wound Care 101: Caring for Developmentally Disabled with Wounds





wound-care-plus-llc



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SECTION M: SKIN CONDITIONS

Intent: The items in this section document the risk, presence, appearance, and change of pressure ulcers. This section also notes other skin ulcers, wounds, or lesions, and documents some treatment categories related to skin injury or avoiding injury. It is important to recognize and evaluate each resident s risk factors and to identify and evaluate all areas at tick of constant pressure. A complete assessment of skin is essential to an effective pressure ulcer prevention and skin treatment program. Be certain to include in the assessment process, a holistic approach. It is imperative to determine the etiology of all wounds and lesions, as this will determine and direct the proper treatment and management of the wound.

CMS is aware or the array of terms used to describe alterations in skin integrity due to pressure. Some of these terms include: pressure ulcer, pressure injury, pressure sore, decubitus ulcer, and bed sore. Acknowledging that clinicians may use and documentation may reflect any of these terms, it is acceptable to code pressure-related skin conditions in Section M if different terminology is recorded in the clinical record, as long as the primary cause of the skin alteration is related to pressure. For example, if the medical record reflects the presence of a Stage 2 pressure injury, it should be coded on the MDS as a Stage 2 pressure ulcer.

M0100: Determination of Pressure Ulcer Risk

M0100. Determination of Pressure Ulcer Risk

Check all that apply

- A. Resident has a stage 1 or greater, a scar over bony prominence, or a non-removable dressing/device
- B. Formal assessment instrument/tool (e.g., Braden, Norton, or other
- C. Clinical assessment
- Z. None of the above

Item Rationale

Health-related Quality of Life

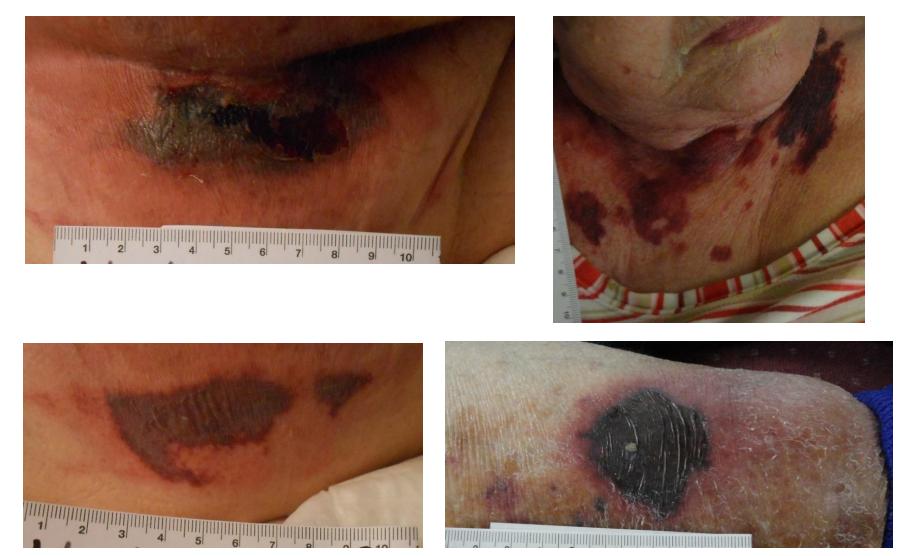
- Pressure ulcers occur when tissue is compressed between a bony prominence and an
 external surface. In addition to pressure, shear force, and friction are important
 contributors to pressure ulcer development.
- The underlying health of a resident's soft tissue affects how much pressure, shear force, or friction is needed to damage tissue. Skin and soft tissue changes associated with aging, illness, small blood vessel disease, and malnutrition increase vulnerability to pressure ulcers.
- Additional external factors, such as excess moisture, and tissue exposure to urine or feces, can increase risk.

Planning for Care

 The care planning process should include efforts to stabilize, reduce, or remove underlying risk factors; to monitor the impact of the interventions; and to modify the interventions as appropriate based on the individualized needs of the resident. Source: CMS's RAI Version 3.0 Manual Page M-1

"It is imperative to determine the etiology of all wounds and lesions, as this will determine and direct the proper treatment and management of all wounds"

Four different etiologies...only one is pressure. Which one is pressure?





Pressure-Unstageable due to DTI



Coumadin Injury



Purpura



Diabetic Wound of the Lower Extremity-Wagner Grade 0



Four different diagnosis. None are pressure.









Sickle Cell Ulcer



Squamous Cell Carcinoma



Pyoderma Gangrenosum



Venous Stasis Ulcer secondary to Chronic Lymphadema

Collagen Vascular Diseases Pyoderma Gangrenosum Sarcoidosis Lesions Typical (Classic) **Digital Sclerosis Discoid Lupus Skin Lesions** Atypical Calciphylaxis Peristomal **Disseminated Granuloma Annulare** Pustular **Arterial Ulcers Diabetic Ulcer** Bullous Venous Insufficiency Ulcers Neuropathic Ulcer Vegetative Surgical Wounds Trauma **Eruptive Xanthomatosis** Malignant Lesions **Skin Tears** Majorlin's Ulcer Acanthosis Nigricans Pemphigoid Sweet's Syndrome **Diabetic Dermopathy** Bullous Necrobiosis Lipoidica Diabeticorum Burns Cicatricial **Bullosis Diabeticorum Necrotizing Infection** Gestationis Pemphigus Vasculitis **Radiation Necrosis Ulcer**

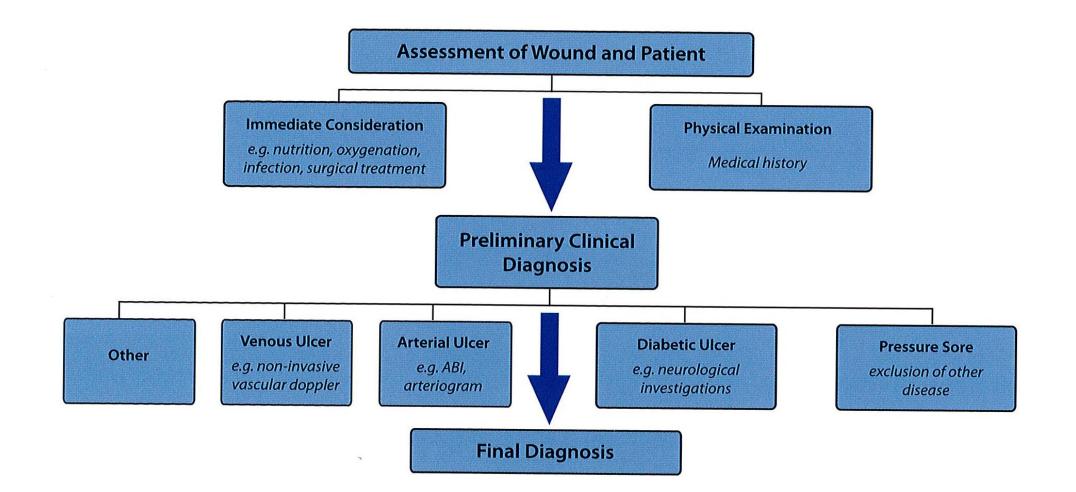
I. Review of a Resident with Non Pressure-Related Skin Ulcer/Wound. Residents may develop various types of skin ulceration. At the time of the assessment and diagnosis of a skin ulcer/wound, the clinician is expected to document the clinical basis (e.g., underlying condition contributing to the ulceration, ulcer edges and wound bed, location, shape, condition of surrounding tissues) which permit differentiating the ulcer type, especially if the ulcer has characteristics consistent with a pressure ulcer, but is determined not to be one. This section differentiates some of the different types of skin ulcers/wounds that are not considered to be pressure ulcers.

Mega Rule Screenshot





At the time of the assessment, clinicians (physicians, advance practice nurses, physician assistants, and certified wound care specialists, etc.) should document the clinical basis (for example, type of skin injury, location, shape, edges and wound bed, condition of surrounding tissues) for any determination that an injury is not pressure-related, especially if the injury has characteristics consistent with a pressure injury, but is determined not to be one.



How long does it take to diagnose a wound or skin issue?

Diagnosed as Pressure and Moisture



Marjolin Ulcer?

Duhring's Disease (aka Dermatitis herpetiformis)



9-28-2016

10-26-2016

How long does it take to diagnose a wound or skin issue?

		CT Scan		
Transcutaneous Oxygen Mor	-	ement	Wedge Biopsy	
Near Infrared S	pectroscopy			
Bacterial Fluorescence	Diagnosis of Exclu	usion	Labo	oratory Draws
		Arteria	l Ultrasound	
Ankle-Brachial Index	X-ray			Tissue Cultures
	Venous	Debridement Laboratory Draws osis of Exclusion Arterial Ultrasound X-ray Tissue Cultures Venous Ultrasound pH Testing Pressure Mapping		
Punch Biopsy			ł	oH Testing
MRI Scan	Arteriogram	Pressure I	Mapping	

Tissue Destruction Classification Systems

Tissue destruction classification systems have been developed to assist describing the depth of penetration of wounds. Some systems are specific to the wound etiology such as the pressure ulcer staging system. Documenting the levels of tissue destruction correctly is critical for communication between clinicians and in some instances, required for reporting to CMS. For instance, CMS regulatory mandates require accurate staging for reporting, reimbursement, and tracking of pressure ulcers in different health care settings.

All Wounds

All wounds, regardless of etiology, can be assessed as either partial or full-thickness.

Classification of Wound by Thickness of Tissue Destruction



Partial-thickness

Extends through the epidermis (first layer of skin), but not through dermis (second layer)



Full-thickness

Extends through epidermis and dermis; may involve subcutaneous tissue, muscles, joint capsule, bone, etc.

Different Classification Scales for Assessing Tissue Destruction

Old System

First degree

Second degree

Third degree

Fourth degree

New System

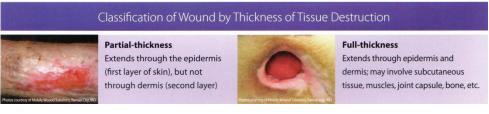
Not classified

Partial-thickness

Full-thickness

Complete burns

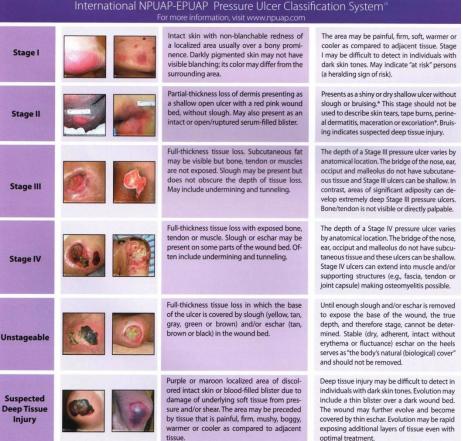
Still in Transition



Assessment by Thickness of Tissue Destruction

		For more information
Stage I		Intact skin with no a localized area usu nence. Darkly pigm visible blanching; its surrounding area.
Stage II		Partial-thickness los a shallow open ulce bed, without slougl intact or open/rupte
Stage III	<u>@</u>	Full-thickness tissue may be visible but are not exposed. Sk does not obscure t May include undern
Stage IV		Full-thickness tissue tendon or muscle. present on some pa ten include underm
Unstageable		Full-thickness tissue of the ulcer is covere gray, green or bro brown or black) in th
	Market Street Street	Purple or maroon

Rev. 2009



Wagner Scale for Diabetic Foot Ulcers"					
Grade	Description				
Grade 0	Pre-ulcerous lesion, healed ulcer, bony deformity				
Grade 1	Superficial ulcer without subcutaneous tissues involvement				
Grade 2	Deep ulcer, penetration through the subcutaneous tissue; may have exposed bone, tendon or ligament or joint capsule				
Grade 3	Deep ulcer with cellulitis, abscess formation, or osteomyelitis				
Grade 4	Localized gangrene of digit				
Grade 5	Extensive gangrene involving whole foot				

1981

Summary

blistered

Fiery red, very painful, but not

Extend through the epidermis and

Healing by regeneration; full function

and appearance should be recovered.

Penetrate the dermis and may involve

subcutaneous tissue. Hair follicles,

sebaceous glands, and sweat glands

are destroyed. Healing occurs through

scar formation and re-epithelialization

Extend into the subcutaneous tissue to

include muscle, fascia or bone. They may generate systemic toxic reactions

or rapidly lead to infection or sepsis.

may penetrate into the dermis.

2010

L1 Hyperemic Lesion Peristomal redness with intact skin



Step 1 Assess and classify the peristomal skin lesion

(L1-LX)





L3 Ulcerative Lesion Open lesion extending into subcutaneous tissue and below: full-thickness skin loss



L4 Ulcerative Lesion Full-thickness skin loss with non-viable, dead tissue (necrotic, fibrinous)





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	Payne-Martin Skin Tear Classification System ^{7,54}
Category I	No tissue loss, may be linear (tears without tissue loss, resembling an incision) or flaps (epidermal flap covers the dermis to within 1 mm of the edge).
Category II	Tissue loss. Scant, partial tissue loss = 25% or less of epidermal flap lost. Moderate to large partial tissue loss = more than 25% of epidermal flap is lost.
Category III	Complete tissue loss; no epidermal flap.

1996

1993		University of Texas Classification System of Diabetic Foot Ulcers				
1000		0			II	Ш
		A	Pre-or post-ulce	rative lesion	Superficial wound, not involving tendon, capsule, or bone	Wound penetrating to tendon or capsule
		В	Pre-or post-ulce	rative lesion	Pre- or post-ulcerative lesion, completely epithelialized with infection	Superficial wound, not involving tendon, capsule, or bone with infection
Skin Tears	kin Tears Rev. 2010		Pre- or post-ulcerative lesion, completely epithelialized with		Superficial wound, not involving tendon, capsule, or bone with ischemia	Wound penetrating to tendon or capsule with ischemia
Star Skin Tear Classification System ^{13,39}			tive lesion, ialized with mia	Superficial wound, not involving tendon, capsule, or bone with infection and ischemia	Wound penetrating to tendon or capsule with infection and ischemia	
	Category 1a A skin tear where the edges can be realigned to the normal anatomical position (without undue stretching) and the skin or flap colour is not pale, dusky or darkened.			ind		Johnma
	Category 1b A skin tear where the edges can be realigned to the normal anatomical					



Category 2a

A skin tear where the edges cannot be realigned to the normal anatomical position and the skin flap colour is not pale, dusky or darkened.

position (without undue stretching) and the skin or flap colour is pale, dusky or darkened.



Category 2b

A skin tear where the edges cannot be realigned to the normal anatomical position and the skin or flap colour is pale, dusky or darkened.



Category 3

A skin tear where the skin flap is completely absent.

Partial Thickness





Stage 2 Pressure Ulcer: Partial-thickness skin loss with exposed dermis

Partial-thickness loss of skin with exposed dermis, presenting as a shallow open ulcer. The wound bed is viable, pink or red, moist, and may also present as an intact or open/ruptured blister. Adipose (fat) is not visible and deeper tissues are not visible. Granulation tissue, slough and eschar are not present. This stage should not be used to describe moisture associated skin damage including incontinence associated dermatitis, intertriginous dermatitis (inflammation of skin folds), medical adhesive related skin injury, or traumatic wounds (skin tears, burns, abrasions).

Stage 3 Pressure Ulcer: Full-thickness skin loss

Full-thickness loss of skin, in which subcutaneous fat may be visible in the ulcer and granulation tissue and epibole (rolled wound edges) are often present. Slough and/or eschar may be visible but does not obscure the depth of tissue loss. The depth of tissue damage varies by anatomical location; areas of significant adiposity can develop deep wounds. Undermining and tunneling may occur. Fascia, muscle, tendon, ligament, cartilage and/or bone are not exposed. If slough or eschar obscures the wound bed, it is an Unstageable PU/PI.

Mega Rule Screenshot



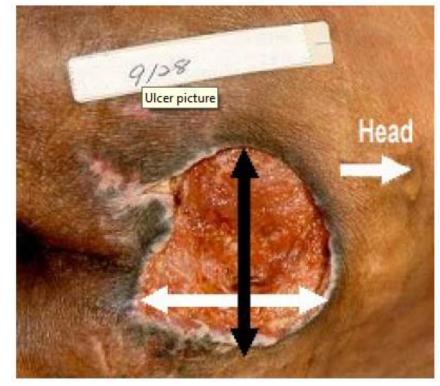






Wound Measurement

- 1. Measurement is based on observation of the Stage 3, Stage 4, or unstageable pressure ulcer due to slough and/or eschar after the dressing and any exudate are removed.
- 2. Use a disposable measuring device or a cotton-tipped applicator.
- 3. Determine longest length (white arrow line) head to toe and greatest width (black arrow line) of each Stage 3, Stage 4, or unstageable pressure ulcer due to slough and/or eschar.
- 4. Measure the longest length of the pressure ulcer. If using a cotton-tipped applicator, mark on the applicator the distance between healthy skin tissue at each margin



and lay the applicator next to a centimeter ruler to determine length.

Standard Tissue Types

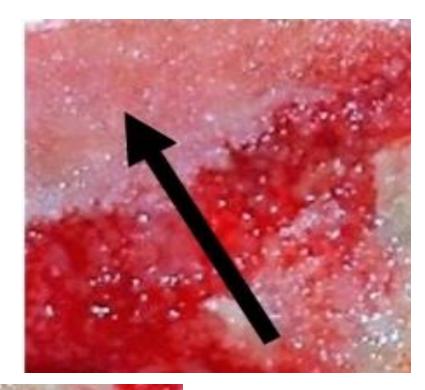
DEFINITIONS

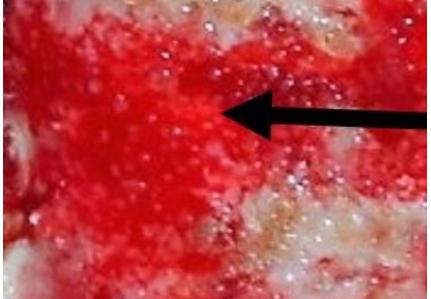
EPITHELIAL TISSUE

New skin that is light pink and shiny (even in persons with darkly pigmented skin). In Stage 2 pressure ulcers, epithelial tissue is seen in the center and edges of the ulcer. In full thickness Stage 3 and 4 pressure ulcers, epithelial tissue advances from the edges of the wound.

GRANULATION TISSUE

Red tissue with "cobblestone" or bumpy appearance, bleeds easily when injured.



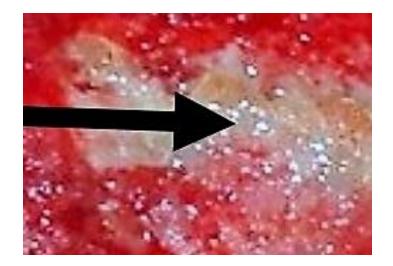


SLOUGH TISSUE

Non-viable yellow, tan, gray, green or brown tissue; usually moist, can be soft, stringy and mucinous in texture. Slough may be adherent to the base of the wound or present in clumps throughout the wound bed.

ESCHAR

Dead or devitalized tissue that is hard or soft in texture; usually black, brown, or tan in color, and may appear scab-like. Eschar is usually firmly adherent to the base of the wound and often the sides/edges of the wound.





Stable Eschar-Do Not Touch

- Dry
- Firmly Adherent
- Eschar Intact
- Peri-wound has no erythema, no fluctuance, no bogginess, no maceration, no warmth, no redness, no induration or swollen tissue

Unstable-Consider having specialist remove

• Drainage

•

- Edges peeling or lifting up
- Eschar does not cover the entire wound
 - Peri-wound may have erythema,
 fluctuance, bogginess, temperature or color
 changes, maceration, warmth or redness,
 induration, or swollen tissue

Stable eschar (i.e., dry, adherent, intact without erythema or fluctuance) on the heels serves as "the body's natural (biological) cover" and should only be removed after careful clinical consideration, including ruling out ischemia, and consultation with the resident's physician, or nurse practitioner, physician assistant, or clinical nurse specialist if allowable under state licensure laws.

Other Tissue Types





Fungating Tumor





Fungating Tumor





Acanthosis Nigricans





Plantar Wart

Wound Exudate

Commons Types

Purulent-opaque, milky and sometimes green, yellow or brown

Sanguineous-reddish, thin and watery

Serosanguineous-clear, pink, thin and watery (blood-tinged)

Serous-clear, amber, thin and watery

Drainage Amount

None-dry wound bed

Scant-wound is moist. Dressing will have no more than a drop or two of drainage present or dressing may be dry

Small-minimal amount of drainage covering less than 25% of the dressing

Moderate-wound is wet and drainage covers 25-75% of the dressing

Large (Copious)-wound has significant fluid that may fill the wound and drainage will cover more than 75% of the bandage. Peri-wound may shows signs of maceration When to Assess

DEFINITION

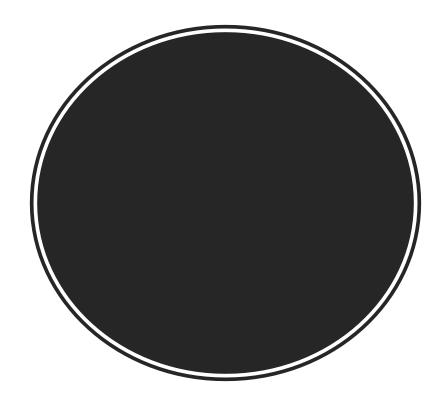
ON ADMISSION

As close to the actual time of admission as possible.



Wound Terminology







Demarcation or Demarcating or Dry Gangrene







Epibole Or Epiboly







Erythema



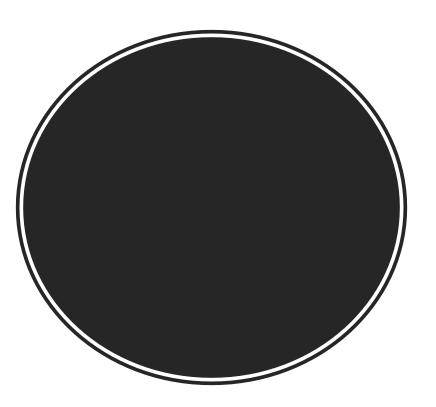




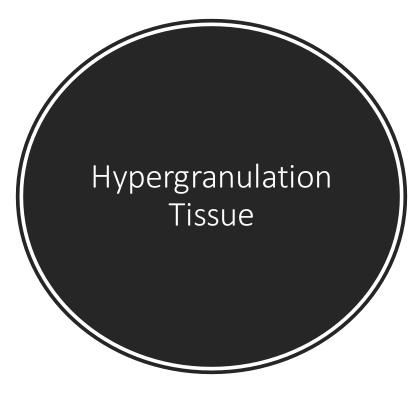


Heralding Sign

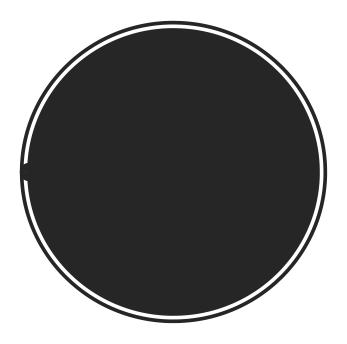




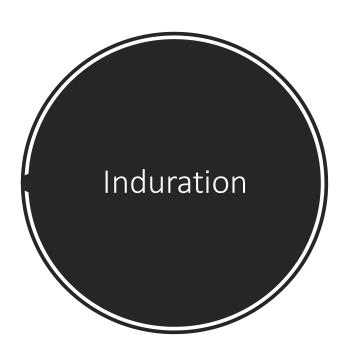




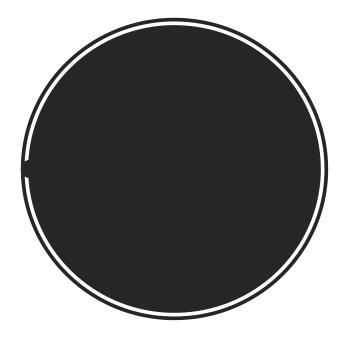


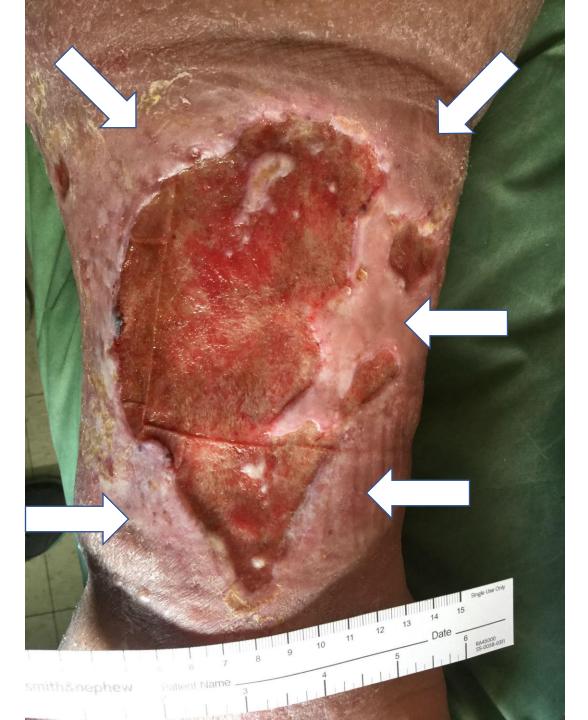


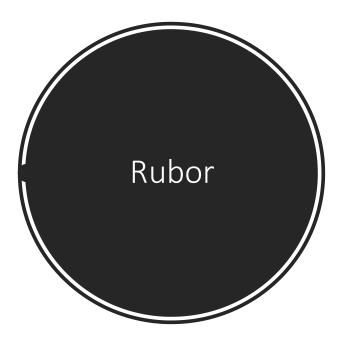


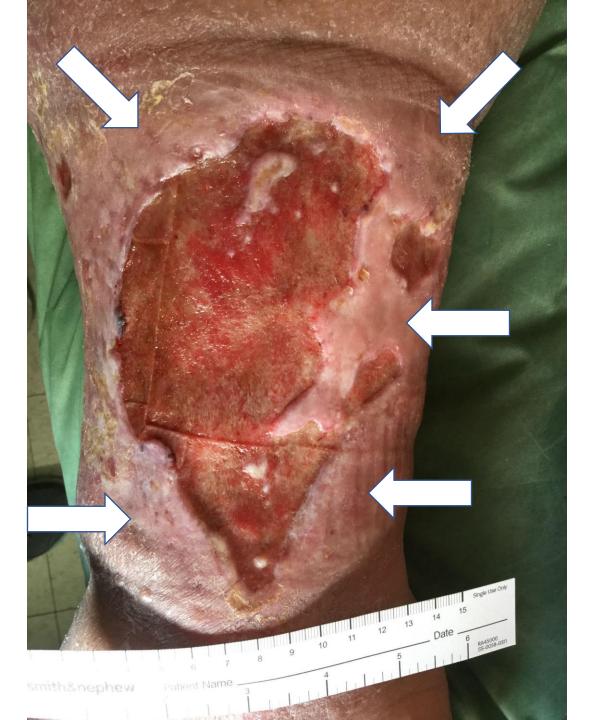












Wound Characteristics Even wound margins Even wound margins Irregular wound margins Gangrene or necrosis Deep wound bed Cellulitis or underlying Deep, pale wound Superficial wound osteomyelitis Ruddy, granular bed Blanched or purpuric Granular tissue tissue periwound tissue present unless PVD is Usually no pain Frequently moderate Severe pain present to heavy exudate Low to moderate Cellulitis drainage Minimal exudate Patient Assessment Firm edema Thin, shinny, dry skin Diminished or absent Hair loss on ankle & sensation in foot Dilated superficial Foot deformities foot veins Thickened toenails Palpable pulses Dry, thin skin Warm foot Evidence of healed Pallor on elevation ulcers and dependent rubor Subcutaneous fat Cyanosis Periwound and leg atrophy hyperpigmintation Decreased Possible dermatitis temperature Absent or diminished pulses

Arterial Ulcers	Diabetic Ulcers	Venous Ulcers
 Wound Characteristics Even wound margins Gangrene or necrosis Deep, pale wound bed Blanched or purpuric periwound tissue Severe pain Cellulitis Minimal exudate 	 Even wound margins Deep wound bed Cellulitis or underlying osteomyelitis Granular tissue present unless PVD is present Low to moderate drainage 	 Irregular wound margins Superficial wound Ruddy, granular tissue Usually no pain Frequently moderate to heavy exudate
 Patient Assessment Thin, shinny, dry skin Hair loss on ankle & foot Thickened toenails Pallor on elevation and dependent rubor Cyanosis Decreased temperature Absent or diminished 	 Diminished or absent sensation in foot Foot deformities Palpable pulses Warm foot Subcutaneous fat atrophy 	 Firm edema Dilated superficial veins Dry, thin skin Evidence of healed ulcers Periwound and leg hyperpigmintation Possible dermatitis

pulses

Medical Adhesive-Related Skin Injury (MARSI)

Similar to medical device related pressure injuries, MARSI describes an etiology of injury. In the case of MARSI the etiology is trauma, not pressure.

The FDA describes medical adhesive tape or adhesive bandages as "a device(s) intended for medical purposes that consists of a strip of fabric material or plastic, coated on one side with adhesive..."

Elderly patients and neonates are at high risk for MARSI. MARSI can be prevented by careful selection of adhesive products, correct application and removal.

The types of skin injuries caused by medical adhesives include:

- Mechanical trauma
 - Epidermal stripping
 - Tension injury or blister
 - Skin tear
- Dermatitis
 - Irritant contact dermatitis
 - Allergic dermatitis
- Other
 - Maceration from trapped moisture
 - Folliculitis

NOTE

There is some overlap in terminology, however, the important thing to

remember is that MARSI should not be counted as pressure injuries.

M0100: Determination of Pressure Ulcer Risk

M0100. Determination of Pressure Ulcer Risk					
↓ Check all that apply					
	A. Resident has a stage 1 or greater, a scar over bony prominence, or a non-removable dressing/device				
	B. Formal assessment instrument/tool (e.g., Braden, Norton, or other)				
	C. Clinical assessment				
	Z. None of the above				



Check B if a formal assessment has been

completed. An example of an established pressure ulcer risk tool is the *Braden Scale* for *Predicting Pressure Sore Risk*^{\odot}. Other tools may be used.

- Check C if the resident's risk for pressure ulcer development is based on clinical assessment. A clinical assessment could include a head-to-toe physical examination of the skin and observation or medical record review of pressure ulcer risk factors. Examples of risk factors include the following:
 - impaired/decreased mobility and decreased functional ability
 - co-morbid conditions, such as end stage renal disease, thyroid disease, or diabetes mellitus;
 - drugs, such as steroids, that may affect wound healing;
 - impaired diffuse or localized blood flow (e.g., generalized atherosclerosis or lower extremity arterial insufficiency);
 - resident refusal of some aspects of care and treatment;
 - cognitive impairment;
 - urinary and fecal incontinence;
 - under nutrition, malnutrition, and hydration deficits; and
 - healed pressure ulcers, especially Stage 3 or 4 which are more likely to have recurrent breakdown.

	RISK FACTORS-If the answer is yes place check in the box. If no leave blank	Date	Date	Date	Date
1	Moribund (Actively dying, imminently terminal)				
2	Impaired/decreased mobility, decreased functional ability				
3	Restraints in place				
4	Comorbid conditions (Does the client have 2 or more chronic diseases or conditions simultaneously such as diabetes, cardiovascular, pulmonary, or renal disease)				
5	Impaired blood flow or diagnosis of atherosclerosis, arterial insufficiency, PVD, chronic edema, smoking or CAD				
6	Refuses or resistant to some aspects of care and/or treatment				
7	Cognitive impairment				
8	Exposure of skin to urinary and/or fecal incontinence, perspiration, drainage, weeping				
9	Poor or reduced meal intake				
10	Poor or reduced fluid intake				
11	Previous healed ulcer and/or open area				
12	At risk for friction or shearing from repositioning or repetitive movements by the client				
13	Admitted with potential for Deep Tissue Injury secondary to preadmission factors like prolonged bed rest, surgery; signs of skin impairment on admission; ambulance transport longer than 1 hour from point of departure to point of arrival				

i		 1	 1
14	Neuropathy and/or decreased sensation to feet and/or lower extremities		
15	Disease or drug therapy that may affect wound healing; including anticoagulant		
	therapy, chemotherapeutic agents, immunosuppressant therapy such as steroids		
16	Medically necessary interventions that may contribute to wound development; such as		
	cast, braces, oxygen tubing, catheter, Head of Bed elevated order		
17	Acute changes in health status		
18	Inpatient/Outpatient hospitalization in the last 90 days		
19	Emergency Room visit in the last 90 days		
20	Current open ulcer		
21	Other (chart in <u>nurses</u> notes)		
	OTHER CLINICAL INFORMATION		
1	Resident Weight: Enter most recent weight		
2	Resident left facility since last report date: Review in nurses notes		
2a	Hospital Admission Date:		
2b	Emergency Room Visit Date:		
2c	Returned from hospital admission during report week. Date of return:		
2d	Other:		
3	(Optional) Braden Score: Please write Braden Score at time of this report		
	Initials		

Arterial Ulcer: An arterial ulcer is ulceration that occurs as the result of arterial occlusive disease when non-pressure related disruption or blockage of the arterial blood flow to an area causes tissue necrosis. Inadequate blood supply to the extremity may initially present as intermittent claudication. Arterial/Ischemic ulcers may be present in individuals with moderate to severe peripheral vascular disease, generalized arteriosclerosis, inflammatory or autoimmune disorders (such as arteritis), or significant vascular disease elsewhere (e.g., stroke or heart attack). The arterial ulcer is characteristically painful, usually occurs in the distal portion of the lower extremity and may be over the ankle or bony areas of the foot (e.g., top of the foot or toe, outside edge of the foot). The wound bed is frequently dry and pale with minimal or no exudate. The affected foot may exhibit: diminished or absent pedal pulse, coolness to touch, decreased pain when hanging down (dependent) or increased pain when elevated, blanching upon elevation, delayed capillary fill time, hair loss on top of the foot and toes, toenail thickening;

Venous or Stasis Ulcer: A venous ulcer (previously known as a stasis ulcer) is an open lesion of the skin and subcutaneous tissue of the lower leg, often occurring in the lower leg around the medial ankle. Venous ulcers are reported to be the most common vascular ulceration and may be difficult to heal, may occur off and on for several years, and may occur after relatively minor trauma. The ulcer may have a moist, granulating wound bed, may be superficial, and may have minimal to copious serous drainage unless the wound is infected. The resident may experience pain that may increase when the foot is in a dependent position, such as when a resident is seated with her or his feet on the floor. *Recent literature implicates venous hypertension as a causative factor. Venous* hypertension may be caused by one (or a combination of) factor(s) including: loss of (or compromised) value function in the vein, partial or complete obstruction of the vein (e.g., deep vein thrombosis, obesity, malignancy), and/or failure of the calf muscle to pump the blood (e.g., paralysis, decreased activity). Venous insufficiency may result in edema and induration, dilated superficial veins, dry scaly crusts, dark pigmented skin in the lower third of the leg, or dermatitis. The pigmentation may appear as darkening skin, tan or purple areas in light skinned residents and dark purple, black or dark brown in dark skinned residents. Cellulitis may be present if the tissue is infected.



HEMOSIDERIN

An intracellular storage form of iron; the granules consist of an ill-defined complex of ferric hydroxides, polysaccharides, and proteins having an iron content of approximately 33% by weight. It appears as a dark yellow-brown pigment.

• **Diabetic Neuropathic Ulcer:** A diabetic neuropathic ulcer requires that the resident be diagnosed with diabetes mellitus and have peripheral neuropathy. The diabetic ulcer characteristically occurs on the foot, e.g., at mid-foot, at the ball of the foot over the metatarsal heads, or on the top of toes with Charcot deformity ; and

Arterial Ulcers

Trophic skin changes (e.g., dry skin, loss of hair growth, muscle atrophy, brittle nails) may also be present. The wound may start with some kind of minor trauma, such as hitting the leg on a wheelchair. The wound does not typically occur over a bony prominence, however, can occur on the tops of the toes. Pressure forces play virtually no role in the development of the ulcer, however, for some residents, pressure may play a part. Ischemia is the major etiology of these ulcers. Lower extremity and foot pulses may be diminished or absent.

Venous Ulcers

The wound may start with some kind of minor trauma, such as hitting the leg on a wheelchair. The wound does not typically occur over a bony prominence, and pressure forces play virtually **no** role in the development of the ulcer.

DEFINITIONS

Definitions are provided to clarify clinical terms related to pressure injuries and their evaluation and treatment.

"Pressure Ulcer/Injury (PU/PI)" refers to localized damage to the skin and/or underlying soft tissue usually over a bony prominence or related to a medical or other device. A pressure injury will present as intact skin and may be painful. A pressure ulcer will present as an open ulcer, the appearance of which will vary depending on the stage and may be painful. The injury occurs as a result of intense and/or prolonged pressure or pressure in combination with shear. The tolerance of soft tissue for pressure and shear may also be affected by skin temperature and moisture, nutrition, perfusion, co-morbidities and condition of the soft tissue.

Friction/Shearing

- *"Friction" is the mechanical force exerted on skin that is dragged across any surface.*
- "Shearing" occurs when layers of skin rub against each other or when the skin remains stationary and the underlying tissue moves and stretches and angulates or tears the underlying capillaries and blood vessels causing tissue damage.

Question 1: Was the ulcer caused by pressure or pressure and shear?

- Is or was patient or the patient's body part immobile?
- Evaluate location of the ulcer. Is it directly from positioning in bed? Chair? Shoe? Device?
- Observe the patient in actual supine, side lying, sitting, and with their devices (splints, braces, etc). Is the specific area of breakdown consistent with applied external force?
- Determine if the patient had a recent period of immobility, including just prior to admission. Consider falls at home, rhabdomylisis, and surgical procedures > 4 hours.

No/unsure

- Rationale: Not all ulcers are from pressure. If the cause was clearly not from pressure, do NOT label as
 pressure.
- **Determine wound type:** Review with the patient's physician, nurse practitioner. In the case that the wound type is unclear, a wound consultation by a Certified Wound Specialist (CWS) may be helpful to determine wound type. Obtain final wound type diagnosis by the patient's physician, nurse practitioner.
- **Document:** In cases where a wound is deemed to be NOT pressure, but has characteristics of pressure, the rationale should be documented.
- **Communicate:** Provide education on wound type, plan of care, prognosis, interventions, treatments, etc. with interdisciplinary team, patient and responsible party

Yes

• **Rationale:** In the case that wound type is deemed pressure, a full evaluation should be completed to determine the extent of the issue, mitigate the pressure cause, determine plan of care and wound type prognosis and communicate effectively with the interdisciplinary team, the patient, and the family.

Dressings and Treatments

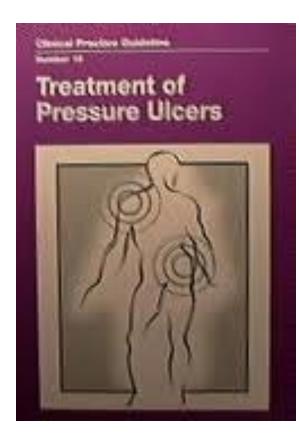
Determination of the need for treatment for a PU/PI is based upon the individual practitioner's clinical judgment, facility protocols, and current professional standards of practice.

Product selection should be based upon the relevance of the specific product to the identified PU/PI(s) characteristics, the treatment goals, and the manufacturer's recommendations for use. Current literature does not indicate significant advantages of any single specific product over another, but does confirm that not all products are appropriate for all PU/PIs. Wound characteristics should be assessed throughout the healing process to assure that the treatments and dressings being used are appropriate to the nature of the wound.

www.woundsource.com

Did you know????

 In 1992 the Agency for Health Care Policy and Research (AHCPR) Guidelines written by the Department of Health and Human Services stated Betadine, Dakin's Solution, and Peroxide should not be used because these products are cytotoxic and not beneficial for wound healing?



(Front page of AHCPR Guidelines Pamphlet)

Why do we care? What do we do about it?

Dressings

- Alginates (35)
- Antimicrobial Dressings (165)
- · Collagens (23)
- · Composites (26)
- Contact Layers (21)
- Foam Dressings (148)
- Gauzes & Non-Wovens (32)
- Gelling Fiber Dressings (9)
- Hydrocolloids (51)
- Hydrogels: Amorphous (32)
- Hydrogels: Impregnated (10)
- · Hydrogels: Sheets (12)
- Impregnated Dressings (32)
- Medical Grade Honey (23)
- · Silicone Gel Sheets (4)
- Specialty Absorptives/Super Absorbents (47)
- Transparent Films (33)
- Wound Fillers (8)

711

http://www.woundsource.com/product-category/dressings



Conditions -

Categories -

All Categories

Biophysical Agents

Practice Accelerator -

-

White Papers

RTICLES WITH A AR AND QUIZ TO CE YOUR WOUND PRACTI

Now

IS

R AND QUIZ TO	Cellular and/or Tissue-Based Products >		
E YOUR WOUND	Compression	F	
	Debridement	×,	
RACTICE	Dressings		Alginates
CODER"	Equipment	F	Antimicrobial Dressings
	Growth Factors	F	Collagens
Alt- N	Moisture Management	F	Composites
DW	Negative Pressure Wound Therapy	F	Contact Layers
1988 -1	Nutritional Management	F	Foam Dressings
	Offloading Devices	►	Gauzes & Non-Wovens
	Positioners & Protectors)	Gelling Fiber Dressings
The World's	Practice Management Software	F	Hydrocolloids
	Pressure Assessment & Monitoring	ŀ	Hydrogels: Amorphous
Product Categories	Skin Care)÷	Hydrogels: Impregnated
> Autologous Delivery Technology	Support Surfaces	ŀ	Hydrogels: Sheets
 Biophysical Agents Cellular and/or Tissue-Based Products 	Tapes & Securement	F	Impregnated Dressings
Wounds	Wound Assessment & Documentation >		Medical Grade Honey
CompressionDebridement	Wound Care - Other	Þ	Silicone Gel Sheets
 Dressings Equipment 	Wound Care Chairs	F	Specialty Absorptives / Super Absorbents
> Growth Factors	Wound Care Diagnostics)	Transparent Films
 Moisture Management Negative Pressure Wound Therapy 	Wound Care Training Tools	F	Wound Fillers

Blog

Webinars

Wounds may be classified as infected if the signs and symptoms of infection are present and/or a wound culture (obtained in accord with accepted standards, such as sterile tissue aspirate, a "quantitative surface swab" using the Levine technique or semi-quantitative swab) contains $100,000 (10^5)$ or greater micro-organisms per gram of tissue. A superficial swab may show the presence of bacteria, but is not a reliable method to identify infection.

Some facilities may use "wet to dry gauze dressings" or irrigation with chemical solutions to remove slough. The use of wet-to-dry dressings or irrigations may be appropriate in limited circumstances, but repeated use may damage healthy granulation tissue in healing ulcers and may lead to excessive bleeding and increased resident pain.

Wet to Dry

- Been around for centuries
- Better products
- Can complicate healing
- Pick your tag





M1200: Skin and Ulcer Treatments (cont.)

- Additional supplementation above the US RDI has not been proven to provide any further benefits for management of skin problems including pressure ulcers. Vitamin and mineral supplementation should only be employed as an intervention for managing skin problems, including pressure ulcers, when nutritional deficiencies are confirmed or suspected through a thorough nutritional assessment (AMDA PU Guideline, page 6). If it is determined that nutritional supplementation, i.e. adding additional protein, calories, or nutrients is warranted, the facility should document the nutrition or hydration factors that are influencing skin problems and/or wound healing and "tailor nutritional supplementation to the individual's intake, degree of under-nutrition, and relative impact of nutrition as a factor overall; and obtain dietary consultation as needed," (AMDA PU Therapy Companion, page 4).
- It is important to remember that additional supplementation is not automatically required for pressure ulcer management. Any interventions should be specifically tailored to the resident's needs, condition, and prognosis (AMDA PU Therapy Companion, page 11).

Based upon the assessment and the resident's clinical condition, choices and identified needs, basic or routine care could include, but is not limited to, interventions to:

- *Redistribute pressure (such as repositioning, protecting and/or offloading heels, etc.);*
- *Minimize exposure to moisture and keep skin clean, especially of fecal contamination;*
- *Provide appropriate, pressure-redistributing, support surfaces;*
- Provide non-irritating surfaces; and
- Maintain or improve nutrition and hydration status, where feasible. Adverse drug reactions related to the resident's drug regimen may worsen risk factors for development of, or for non-healing PU/PIs (for example, by causing lethargy or anorexia or creating/increasing confusion) and should be identified and addressed. These interventions should be incorporated into the plan of care and revised as the condition of the resident indicates.

DEFINITIONS

PRESSURE ULCER/ INJURY RISK FACTOR

Examples of risk factors include immobility and decreased functional ability; co-morbid conditions such as end-stage renal disease, thyroid disease, or diabetes; drugs such as steroids; impaired diffuse or localized blood flow; resident refusal of care and treatment; cognitive impairment; exposure of skin to urinary and fecal incontinence; *microclimate*, malnutrition, and hydration deficits; and a healed ulcer.











3M[™] Cavilon[™] Advanced Skin Protectant is an ultra-thin transparent barrier that protects vulnerable skin from caustic, corrosive body fluids without impacting visualization of the skin. Waterproof, flexible, irritant-proof.

Benefits

• Combines a unique polymer system with cyanoacrylate to create a highly durable, ultra-thin, transparent barrier with elastomeric properties

• Has been shown to significantly improve severe cases of IAD - even in the presence of continued incontinence

- Prevents skin injury and maintains skin integrity
- Reduces the pain associated with managing IAD
- Reduces overall time spent on the prevention and management of IAD and other types of skin damage
- Supports infection-control initiatives
- Lasts up to seven days
- Attaches to wet, weepy or damaged skin

Arjo's Skin IQ[®] Microclimate Manager (MCM) is a fluid resistant, vapor permeable, single patient use mattress cover for use with a pressure redistribution surface to help prevent and treat Stage I-IV pressure ulcers. Powered by Negative Airflow Technology (NAT), *Skin IQ MCM* continually draws away excess moisture from the skin/surface interface and helps control skin temperature.

Effective and comfortable for patients

- With an outstanding Moisture Vapor Transfer Rate (MVTR) of 130 (g/m2)/hr⁽¹⁾, Skin IQ is more effective in microclimate management than conventional Low Air Loss (LAL) surfaces, which have an average MVTR of 97.7 (g/m2)/hr⁽²⁾
- Helps reduce shear and friction⁽³⁾
- Helps improve patient comfort: in a study of 16 healthy participants using *Skin IQ MCM* on a pressure redistribution surface, it significantly improved the level of comfort⁽²⁾



Determine repositioning frequency with consideration to the individual's:

- Level of activity and mobility,
- General medical condition,
- Overall treatment objectives,
- Skin condition, and
- Comfort.

The resident's skin condition and general comfort should be regularly assessed. The efficacy of repositioning must be monitored and revisions to the care plan considered, if the individual is not responding as expected to the repositioning interventions.







Wound #1 Buttock Before Picture 10/11/17 Wound #1 Buttock After Picture 11/1/17

Case presented as a rash ongoing for the last year or more. Previous treatments included moisture barrier, antifungal, various brief trials, good pericare without significant changes. A punch biopsy was performed in October 2017. Results were negative for malignancy and positive for telangiectasia with sparse perivascular mononuclear cell infiltrate. Patient was subsequently started on estrogen cream and antifungal ointment simultaneously. Rash resolved within 3 weeks and patient was discharged off caseload with 100% resolution of symptoms and complaints.











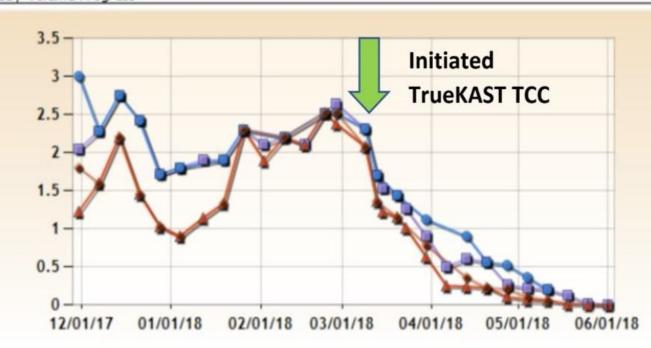






Case is ongoing treatment of a Diabetic wound of the right heel that has been present for 2 years. Wound Care Plus, LLC started treating the patient in November 2017. Calcium alginate was the treatment recommended on the first visit. Xray prior to application of total contact cast was negative for osteomyelitis and positive for a heel spur. ABI 0.9. Client propels self in wheelchair utilizing right heel for mobility. Patient is obese with poor nutritional intake. Physical therapy consulted for strength and mobility training. At time of application of cast, treatment was Santyl and calcium alginate covered with an ABD pad. Quality of tissue was deteriorating.

Area / Volume Progress



Pre Debridement Area
 Pre Debridement Volume
 Post Debridement Area
 Post Debridement Volume



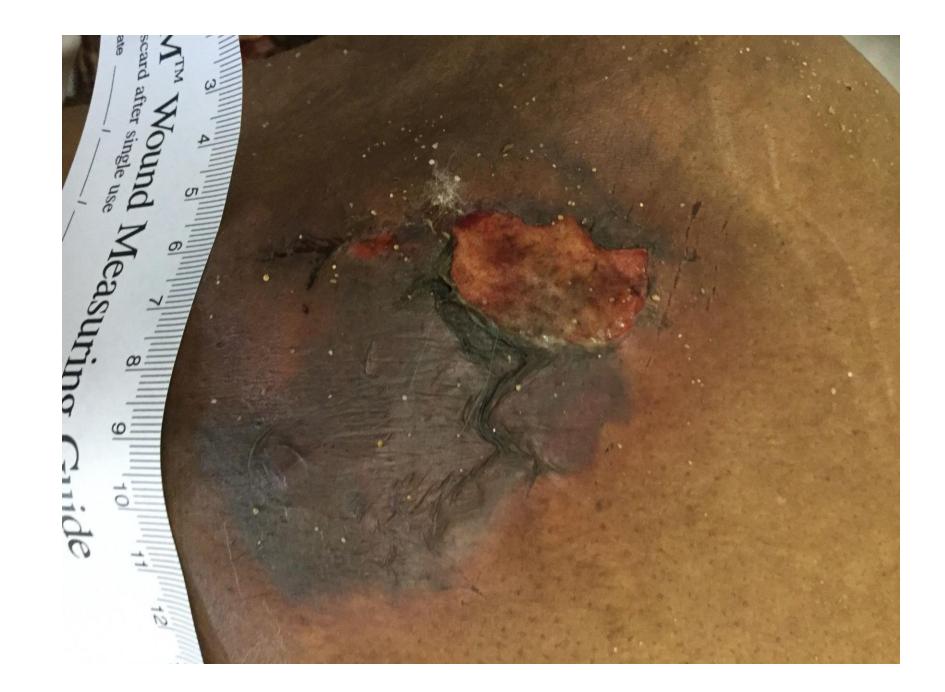














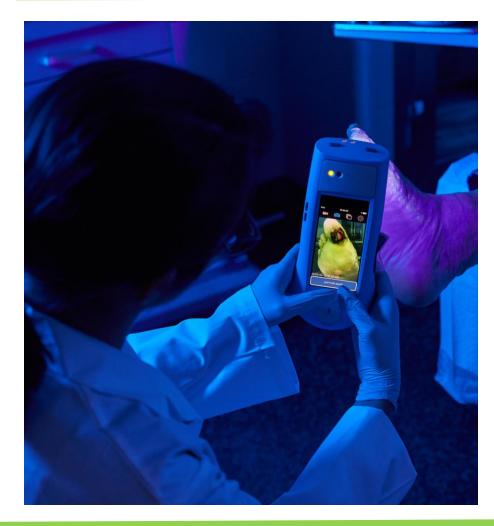
Ulceration of Abdominal Striae Distensae





- 1. Why is the wound there?
- 2. How do you know?
- 3. What's keeping it from healing?
- 4. What are you doing about it?

Enhancing Wound Assessment with MolecuLight *i*:X



- Point-of-care fluorescence imaging device to:
 - 1. Detect bacteria (>10⁴ CFU/g) in wounds^{*}
 - 2. Digitally measure wound area
- Supports point-of-care wound assessment and documentation
- Handheld, portable, touch-screen
- No contrast agents and no patient contact required

*when used with clinical assessment





What does the MolecuLight *i*:X show?

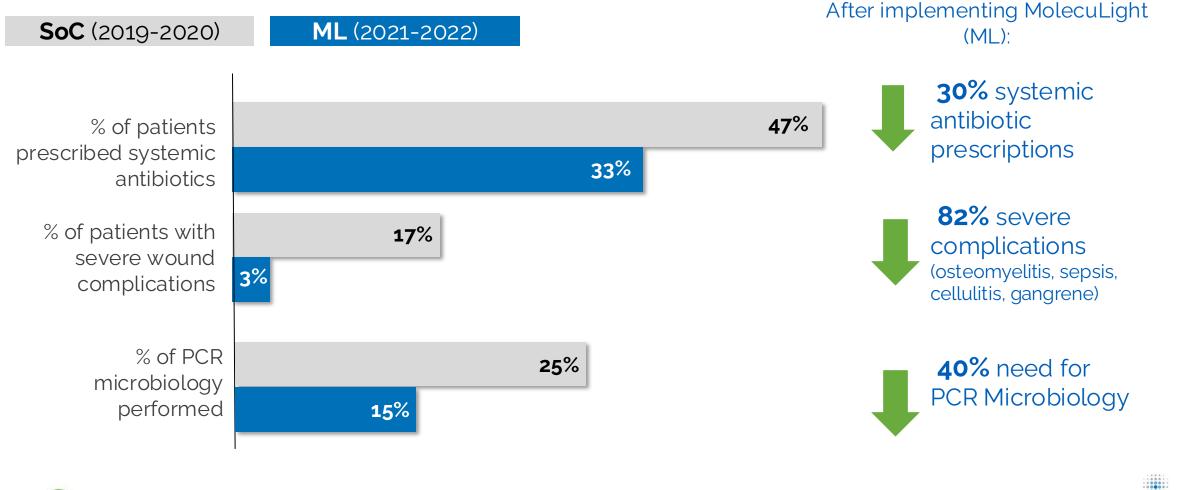






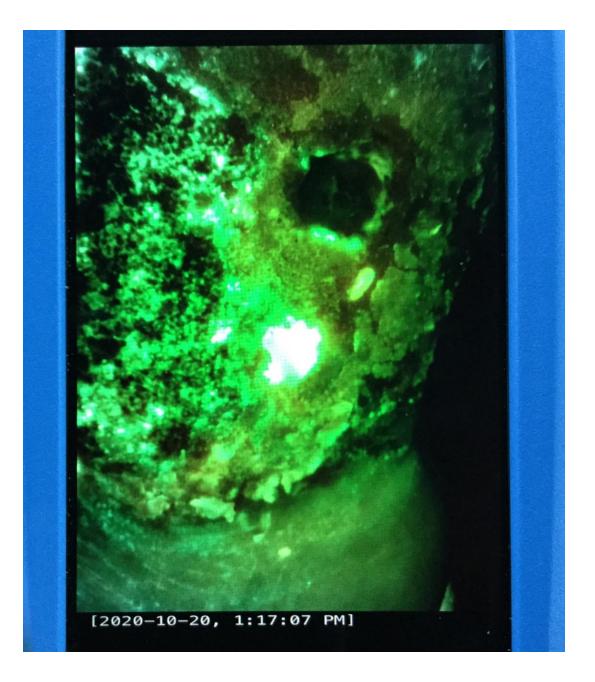
Results

RWE of improved outcomes in Medicare beneficiaries (SNF & LTC)



MolecuLight



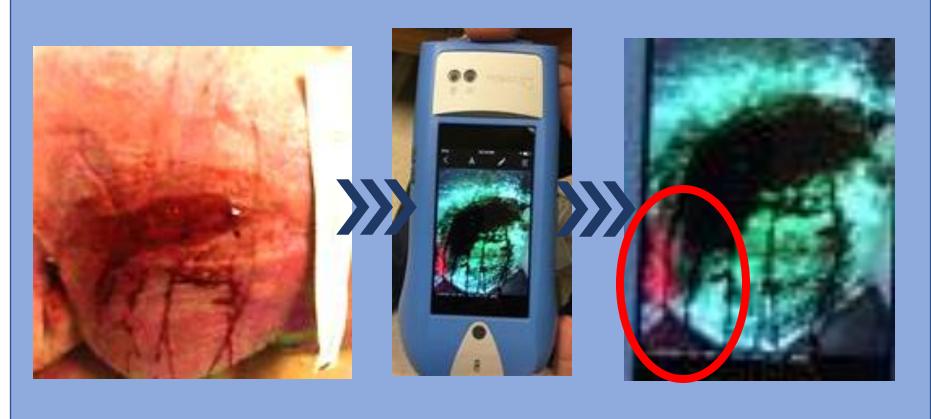


Real World Case Examples

RWE of improved outcomes in Medicare beneficiaries (SNF & LTC)

The value of reducing clinical uncertainty through the localization of bacterial loads

- Targeted treatment
- Accurate tissue sampling site location
- Confirm the need for intervention (i.e debridement or antibiotics)

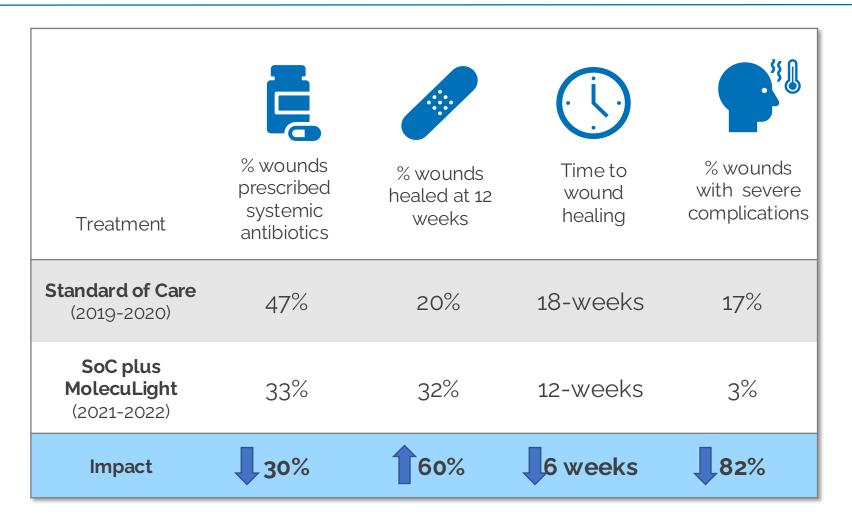






Summary of Findings

RWE of improved outcomes in Medicare beneficiaries (SNF & LTC)







Targeted Debridement, Targeted Cultures

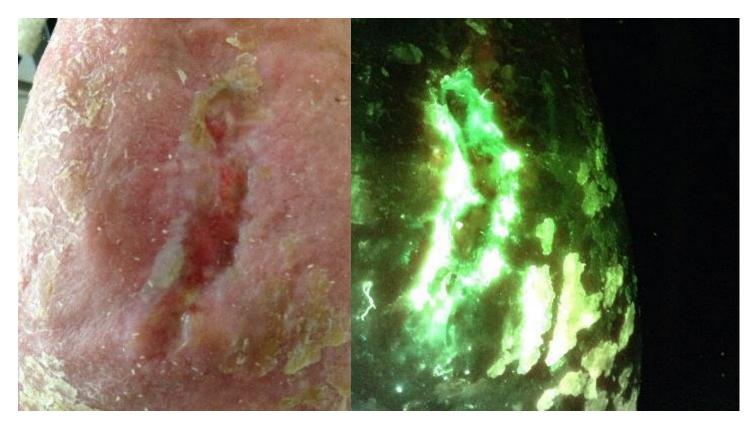








Fluorescence Imaging: Reasonable and Medically Necessary



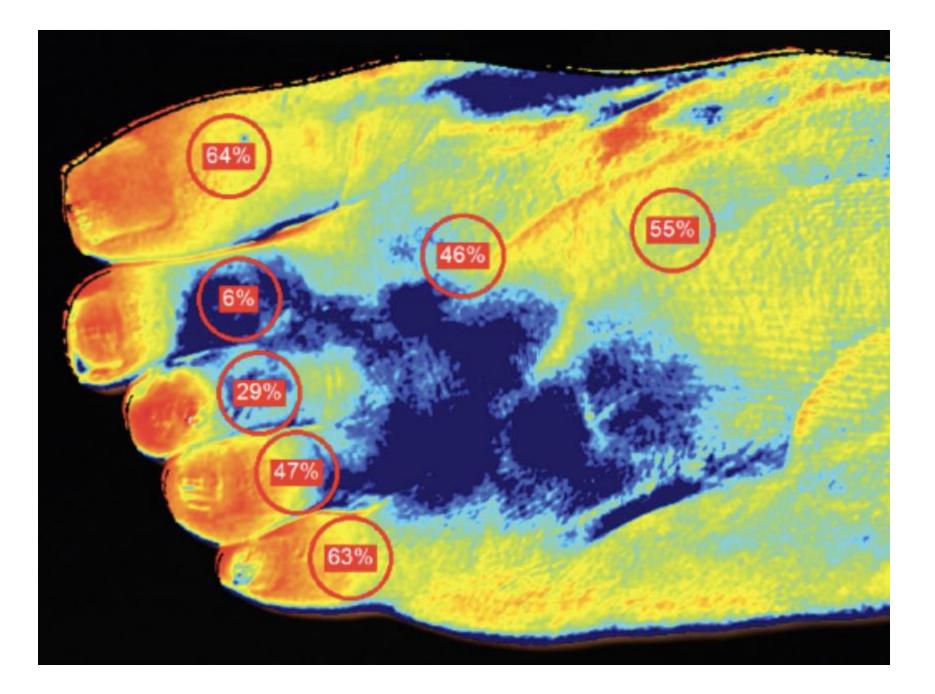
Wound biopsy indicated bacterial loads >10⁶ CFU/g (after more than 3 days delay)

Peer reviewed publications on fluorescence imaging procedure from over 1,200 patients across all patient care settings consistently reports:

- 300-400% increase in detection of high bacterial burden^{1,2}
- >95% accuracy³⁻⁵
- 2.4-fold increase in wound healing rate⁶
- Less amputation, reduced antimicrobial use^{6,7}

1. Serena et al. J Wound Care, 2019; 2. Le et al. Adv. Wound Care, 2021; 3. Rennie et al. J Wound Care 2017; 4. Hurley et al. J Wound Care, 2019; 5. Raizman et al. Diagnostics, 2021; 6. Price N, Diagnostics, 2020; 7. Hill & Woo, Wounds, 2020

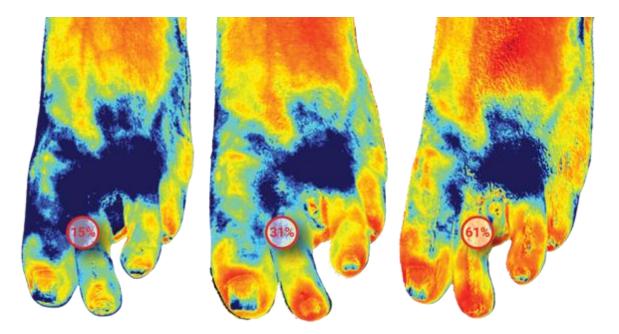




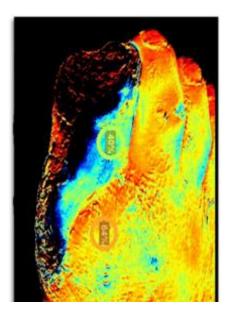
When Time is Tissue

Snapshot_{NIR} helps to expedite vascular referral and assess treatment effectiveness

- Triage patients earlier in the care stream
- Provides point-of-care actionable data to assess tissue response to therapy
- Understanding tissue viability in the wound and the peri-wound can enhance wound bed preparation
- Documents medical necessity and outcomes to help support third-party payment



Surgical Decisions See what lies beneath



- History of arterial disease
- Developed a gangrenous
 great toe



- Great toe amputation
- Kent Image shows flap at risk while clinical image appears normal



 Initial amputation demonstrated continued failure, which led to secondary amputation

How would you intervene and practice limb preservation?

got telehealth? Wound Care Plus



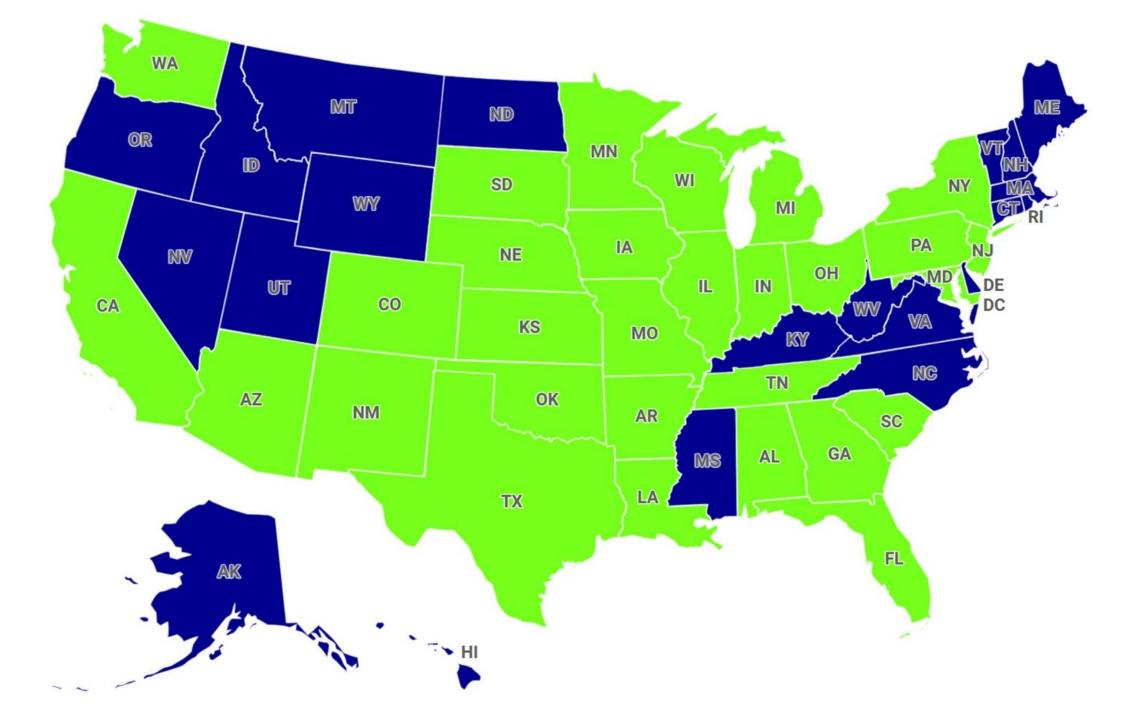
In Person Rounds



We can treat all payor sources (Med A Skilled, Medicaid, Medicaid pending, Private Insurance, Hospice, etc). We can treat all kinds of skin and wound issues (pressure, arterial, diabetic, pemphigoid, rashes, stalled skin tears, blisters, etc).







"Do the best you can until you know better. Then when you know better, do better." - Maya Angelou

Questions?

Martha R Kelso, RN, CHWS, DAPWCA, LNC, HBOT Chief Executive Officer



