Incontinence Associated Dermatitis
or
Pressure Injury?

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**Introduction**

Due to similar locations and presentations, moisture lesions are often misclassified as pressure ulcers\(^1\). Training programs which help to teach us to differentiate between the two have been shown to be beneficial in producing more accurate diagnoses\(^2\). The two aetiologies each have a unique treatment focus so we need to understand the differences between them. Differences are summarized in Appendix A.

**Incontinence Associated Dermatitis (IAD)**

**Definition**

IAD is inflammation of the skin of the perianal or genital areas, buttocks, or upper thighs due to prolonged contact with faeces, urine or sweat\(^3\). Depending on the severity it may or may not be associated with loss of superficial skin layers and/or secondary infections\(^4\). The presence of infection, often by opportunistic fungi, further increases morbidity\(^5\).

In individuals with darker-toned skin IAD may present as areas of hypo- or hyper-pigmentation\(^5\).

**Incidence**

Reported prevalence rates vary greatly from 5% to almost 50% in the literature. The lower numbers are reported from nursing homes, mid range numbers are from hospitals (as a percentage of incontinent patients) and the highest rates are seen in acute geriatric care facilities\(^4,6\). One study in an intensive care unit (ICU) found that 36% of the patients in the study developed IAD within an average of 4 days. The patients in this study had faecal incontinence and diminished cognitive awareness; these characteristics were determined to be significant independent risk factors for IAD development\(^5\).

As the population ages IAD is likely to become more of a problem. It is estimated that 31% of older women and 23% of older men experience urinary incontinence. 12% of older people also experience faecal incontinence\(^3\).

**Pathophysiology**

Components contributing to IAD are\(^6\):

<table>
<thead>
<tr>
<th>Component</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Impairments in tissue tolerance</td>
<td>This includes but is not limited to aging skin, reduced skin perfusion (as might be related to oedema secondary to congestive cardiac failure), and presence of previous scar tissue.</td>
</tr>
<tr>
<td>Problems of the perineal environment</td>
<td>Incontinence, excessive perspiration</td>
</tr>
<tr>
<td>Altered toileting ability from daily use of restraints</td>
<td>Cognitive or physical ability, wheelchair bound, restraints</td>
</tr>
</tbody>
</table>

Prolonged exposure of the skin to moisture leads to maceration, which weakens the epidermis. Friction (such as that from pulling against bedding, clothing or incontinence aids during repositioning) will have a greater impact on this weakened skin allowing it to shear away\(^3\). Another source of friction is that created by skin folds\(^4\).

The protective barrier of the epidermis relies on its acidic nature. Ammonia (formed when urease-producing bacteria in urine or faeces split urea) is alkaline. When urine or faeces is in prolonged contact with the skin, the pH of the skin increases, reducing its ability to...
combat infection\(^{(3)}\). In addition, faeces contain coliform bacteria and digestive enzymes that encourage greater skin irritation\(^{(4)}\).

How do we treat it?
The main aim when treating IAD is to eliminate the cause - prolonged moisture on the skin. Through cleaning and the use of barrier protection we will also try to simulate the normal function of the skin until it has had time to repair itself.

**Contain or divert the fluid.** Absorbent pads are the mainstay of incontinence management and they are remarkably sophisticated in their absorbency capacity and also their ability to draw moisture away from the skin. However, it has been shown that even these products can actually increased tissue interface pressures when soaked, even when used in conjunction with pressure-reducing or relieving support surfaces\(^{(6)}\). Therefore they should be changed often and in accordance with manufacturer’s instructions. They are also not able to draw away any solid matter that may be present in faecal incontinence. A study into the use of perianal pouches for faecal incontinence showed a reduced incidence of IAD and delayed onset compared to patients managed in adult containment briefs\(^{(4)}\).

**Clean the skin.** Cleaning is also carried out regularly on all patients; incontinent patients tend to require more frequent perineal washes. Unfortunately, most skin cleansers have a pH of around 9, which is quite basic, and their frequent use in combination with course washcloths can increase stratum corneum swelling and alter lipid rigidity, stripping the skin of its natural barrier. Cleansing should involve a product whose pH range reflects the acid mantle of healthy skin (pH between 5.4 and 5.9)\(^{(3, 6, 7)}\).

**Provide a Barrier.** A barrier is thought to assist in reducing the contact between the excess fluid and the skin. Depending on the barrier it may also help to reduce friction. There is some evidence that the use of a barrier can help to prevent and/or treat IAD\(^{(3)}\). In a trial involving a washcloth that was impregnated with a barrier product only 8% of patients developed IAD as opposed to 27% of patients in the control group (n=141, \(P=0.003\))\(^{(6)}\). One popular barrier and cleanser washcloth combination has been shown to improve sin integrity in just a couple of days, not only in the groin but in other skin fold areas as well\(^{(7)}\).

**Moisturize the skin.** Moisturising is known to be beneficial in protecting skin. It repairs the skin barrier, retains and increases water content, reduces transepidermal water vapour loss, and restores the lipid barrier by replacing the intracellular lipids\(^{(6)}\). The presence of a moisturizer in the cleaning routine also appears to reduce IAD. In the ICU study, 10% of patients receiving the moisturizing cleanser developed IAD as opposed to 56% in the control group (n=45). This study also commented on the need for the cleansing routine to follow the patient to the ward post ICU, as complete healing had not been achieved in ICU\(^{(5)}\).
There are many products on the market that provide barrier protection. There tend to be three base components, each of which have different strengths and weaknesses in relation to protection from irritation and ability to hydrate:\(^2\):

<table>
<thead>
<tr>
<th>Product</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Petrolatum</td>
<td>Soft white paraffin, good protection against irritants and maceration, some skin hydration</td>
</tr>
<tr>
<td>Dimethicone</td>
<td>Cavilon, varied protection against irritants, good skin hydration</td>
</tr>
<tr>
<td>Zinc oxide</td>
<td>Sudocrem, Amolin, good protection against irritants, poor skin hydration</td>
</tr>
</tbody>
</table>

**Is there an association between IAD and PI?**

A systematic review and meta-analysis by Beekman et al (2014)\(^8\) looked at research that included urinary incontinence, faecal incontinence, double incontinence and moisture/microenvironment and the incidence and development of pressure injuries. It showed varying strengths to the relationship between the two but generally indicated that there is a clear association that if you have IAD you are at a greater risk of developing a pressure injury. One of the articles cited in the review created interesting food for thought. It stated that in nursing homes with high rates of pressure injuries there were high rates of faecal incontinence, but in nursing homes with low rates of pressure injuries there were still high rates of faecally incontinent residents. So does this mean if we are better at managing incontinence we will we be better at managing pressure injuries, or vice versa?

**Pressure Injury (PI)**

**Definition**

A pressure injury is a localized injury to the skin and/or underlying tissue, usually over a bony prominence, as a result of pressure or pressure in combination with shear. A number of contributing or confounding factors including, but not limited to, friction, microclimate, moisture, and nutrition are also associated with pressure ulcers; the significance of these factors has yet to be elucidated\(^1, 9, 10\).

**Incidence**

Reports from the UK indicate an 18% prevalence rate, USA reports 15%. These figures tend to be higher in palliative care/in home hospice, in patients with spinal cord injuries, and in critical care units. Associated with the incidence of PI are longer hospital lengths of stay and higher readmission rates. Recent European cost-models have indicated that the total costs of PIs may consume between 1% in the Netherlands and 4% in the United Kingdom of health care expenditure. They also have a substantial impact of quality of life for the patients and their carers\(^9\).

Sixty thousand patients die each year from complications associated with PIs with an estimated cost of $11 billion per year to treat them (American data). PIs can delay discharge, expose the patient to potential serious infections, cause pain related to the ulcer and its treatment and is a poor indicator of mortality in an acute care setting\(^11\). PIs are considered to be mostly preventable. Recently in Queensland a system has been introduced which penalizes hospitals for causing these injuries to patients. For a stage 3 or stage 4 PIs acquired in the hospital, the hospital will be fined $30k or $50k respectively.
Pathophysiology
Tissue loading is the defining characteristic of pressure injury formation. Pressure is increased in tissues that are positioned between a bony prominence and a support surface. Research has demonstrated that both magnitude of pressure and duration impact on pressure injury formation\(^{(12)}\). Unrelieved pressure disrupts blood supply to the capillary network, impeding blood flow and depriving tissues of oxygen and nutrients. The most common sites for pressure injuries are the sacrum, heels, ischial tuberosities, greater trochanters, and lateral malleoli\(^{(13)}\). Pressure is relieved and circulation restored by frequently turning and shifting weight distribution as well as with the use of dynamic surfaces that actively redistribute pressure on the body surfaces\(^{(12)}\).

How do we treat it?
The recommended treatment of pressure injuries is multi-faceted, each part addressing the underlying cause or a contributing factor. The main cause of a pressure injury is pressure, so the main focus should also be on pressure removal/reduction. The flowchart of recommended treatment from the Pan Pacific Guidelines is in Appendix B.

Implement skin protection strategies. Skin protection is key in the prevention of PI’s. Intrinsic factors such as nutritional and hydration status (see “Refer to Dietician”) and external factors such as shear, friction and moisture, all need to be managed. The Pan Pacific Guidelines recommend\(^{(10)}\):
- Protecting the skin from moisture through an individualised continence management plan
- Use a pH appropriate skin cleanser (avoid the use of alkaline soaps and cleansers) and dry thoroughly to protect the skin from excess moisture
- Use water-based skin emollients to maintain skin hydration (applying a moisturiser contributes to the maintenance of the healthy skin)
- Consider applying a topical barrier preparation to the peri-wound skin to protect it from exudate
- Minimise shear and friction through appropriate manual handling

Refer to Dietician. There is some research to support the use of additional nutritional support for the prevention and management of PIs where appropriate nutritional intake may be lacking. Oral nutrition support (ONS) includes supplements taken orally in addition to the patient’s regular diet. The most common forms of ONS considered for patients with PIs include\(^{(10)}\):
- high protein supplements,
- disease-specific supplements,
- vitamin or multi-vitamin supplements, and
- arginine containing supplements.

Use appropriate support surfaces, appropriately. A support surface is a surface on which the patient is placed to manage pressure load, shear, friction and microclimate. This includes bed, trolley and operating table mattresses, integrated bed systems, and seat cushions. Support surfaces are designed to reduce interface pressure through increasing the body surface area or alternating the area of the body in contact with the support surface. It is important to note that the use of a support surface does not negate the repositioning requirements of a patient with a PI\(^{(10)}\).

Make use of your available resources such as the PIP CNC, your Occupational Therapists and CERU when selecting an appropriate support surface for the severity of the PI and the
patient’s general condition. Once the patient is using a support surface continue to monitor its effectiveness and consider trying something else if the patient’s skin condition is deteriorating and/or showing no signs of improvement. One example is that of the ROHO cushion where the patient puts the cushion behind their back instead of sitting on it.

**Regular repositioning.** Sustained pressure to areas of the body causes soft tissue injury. In normal circumstances, pain resulting from sustained injury prompts a person to change position. In patients who are unable to reposition themselves due to physical limitations, and in patients with reduced sensory perception and impaired ability to detect pain, failure to reposition is a significant risk factor for PI s. Regular repositioning is an essential component of PI prevention\(^{(10)}\).

**Tips for positioning in bed\(^{(10)}\):**
- Whenever the patient is repositioned assess the patient’s skin condition and general comfort and reconsider frequency and method of positioning if the patient is not responding as expected.
- When repositioning the patient reduce friction and shear forces through use of repositioning or transfer aids.
- Where possible, avoid positioning the patient on bony prominences (including heels) with existing erythema.
- Ensure heels are free of the bed surface and inspect the skin of heels frequently.
- If sitting when head-of-bed elevation is required, use aids such as pillows that support the upper body to reduce additional pressure on the sacrum and coccyx.
- Before raising the head-of-bed, move the patient up the bed and raise the knees. This assists in avoiding shear from the patient slipping down the bed.
- Consider more frequent, smaller shifts in position for patients who cannot tolerate frequent and/or major changes in body position.
- Provide transfer assistance devices (e.g. overhead handle) to promote independent patient transferring and reduce shear forces and friction.

**Tips for positioning in chairs/wheelchairs\(^{(10)}\):**
- Position a seated patient in a posture that minimises pressure, friction and shear forces and maintains their usual range of activity.
- When seated in non-reclining chairs ensure the patient’s lower limbs are supported in optimal alignment (e.g. 90° at hip, knee and foot) within the patient’s range of movement to minimise pressure under ischial tuberosities.
- Consider adjusting the seat height and depth to improve supported body positioning. All patients should have appropriate seat to floor height to reduce potential for shear and friction.

**Correctly classify the PI.** Use NPUAP/EPUAP pressure injury classification system\(^{(10, 14)}\):
Category/Stage I: Non-blanchable redness of intact skin
Intact skin with non-blanchable erythema of a localized area usually over a bony prominence. Discoloration of the skin, warmth, edema, hardness or pain may also be present. Darkly pigmented skin may not have visible blanching. Further description: The area may be painful, firm, soft, warmer or cooler as compared to adjacent tissue. Category/Stage I may be difficult to detect in individuals with dark skin tones. May indicate “at risk” persons.

Category/Stage II: Partial thickness skin loss or blister
Partial thickness loss of dermis presenting as a shallow open ulcer with a red pink wound bed, without slough. May also present as an intact or open/ruptured serum- filled or serosanguinous filled blister. Further description: Presents as a shiny or dry shallow ulcer without slough or bruising. This category/stage should not be used to describe skin tears, tape burns, incontinence associated dermatitis, maceration or excoriation.

Category/Stage III: Full thickness skin loss (fat visible)
Full thickness tissue loss. Subcutaneous fat may be visible but bone, tendon or muscle are not exposed. Some slough may be present. May include undermining and tunneling. **Further description:** The depth of a Category/Stage III pressure ulcer varies by anatomical location. The bridge of the nose, ear, occiput and malleolus do not have (adipose) subcutaneous tissue and Category/Stage III ulcers can be shallow. In contrast, areas of significant adiposity can develop extremely deep Category/Stage III pressure ulcers. Bone/tendon is not visible or directly palpable.

Category/Stage IV: Full thickness tissue loss (muscle/bone visible)
Full thickness tissue loss with exposed bone, tendon or muscle. Slough or eschar may be present. Often include undermining and tunneling. **Further description:** The depth of a Category/Stage IV pressure ulcer varies by anatomical location. The bridge of the nose, ear, occiput and malleolus do not have (adipose) subcutaneous tissue and these ulcers can be shallow. Category/Stage IV ulcers can extend into muscle and/or supporting structures (e.g., fascia, tendon or joint capsule) making osteomyelitis or osteitis likely to occur. Exposed bone/muscle is visible or directly palpable.

Unstageable/ Unclassified: Full thickness skin or tissue loss – depth unknown
Full thickness tissue loss in which actual depth of the ulcer is completely obscured by slough (yellow, tan, gray, green or brown) and/or eschar (tan, brown or black) in the wound bed. **Further description:** Until enough slough and/or eschar are removed to expose the base of the wound, the true depth cannot be determined; but it will be either a Category/Stage III or IV. Stable (dry, adherent, intact without erythema or fluctuance) eschar on the heels serves as “the body’s natural (biological) cover” and should not be removed.
**Suspected Deep Tissue Injury-depth unknown**

Purple or maroon localized area of discolored intact skin or blood-filled blister due to damage of underlying soft tissue from pressure and/or shear. **Further description:** The area may be preceded by tissue that is painful, firm, mushy, boggy, warmer or cooler as compared to adjacent tissue. Deep tissue injury may be difficult to detect in individuals with dark skin tones. Evolution may include a thin blister over a dark wound bed. The wound may further evolve and become covered by thin eschar. Evolution may be rapid exposing additional layers of tissue even with treatment.

**Debride the wound as indicated.** Debridement is commonly performed on wounds to remove non-viable or infected tissue and debris in order to prepare the wound bed to receive therapeutic healing products (wound bed preparation). The aim is to maximise the healing process. Non-viable tissue can prolong the healing process by increasing inflammation, levels of bacteria and toxins, and inhibiting re-epithelialisation. Non-viable tissue generally presents as moist, yellow, green or grey in colour and over time becomes dry black or brown eschar. The most commonly used methods of debridement are:

- surgical sharp,
- conservative sharp,
- autolytic,
- enzymatic,
- larval,
- mechanical

Appendix C compares a list of debridement options.

When debridement is indicated, select the method of debridement with consideration to:

- the patient’s condition (including pain, vascular condition, and bleeding risk - a vascular assessment should be conducted prior to debriding any lower extremity PI),
- comfort (debridement is often a painful intervention, conduct a pain assessment and provide appropriate pain relief before debriding a wound)
- type, quantity and location of non-viable tissue;
- goals of care;
- patient preferences;
- health professional training and experience; and
- availability of resources.

Surgical debridement is appropriate when there is an urgent need to remove non-viable tissue (e.g. advancing cellulitis, sepsis, pain, exudate or malodour). Conservative sharp wound debridement should only be performed by health professionals with appropriate training.
**Treat infection.** Antimicrobial therapy includes topical agents such as cadexomer iodine, silver, honey and other topical antimicrobials, as well as systemic antibiotics. All products should be used following comprehensive assessment and in accordance with the licensing authority endorsement and the manufacturers’ directions. Although all chronic wounds are contaminated or colonised, not all are infected. Signs of local infection in a pressure injury include\(^{(10)}\):

- new wound breakdown/increased wound size,
- erythema localised to the peri-ulcer tissue,
- increased amount of exudate,
- increase in viscosity or purulence of exudate,
- increased or unexplained pain,
- oedema of peri-ulcer tissue,
- increased peri-ulcer tissue temperature,
- malodour; and
- tracking, bridging or pocketing within the tissue or probing to the bone.

Appendix D compares a list of common antibacterial compounds.

Antibiotic resistance is a significant concern due to the over use or inappropriate use of antibiotic therapy. Patients should be advised to complete their antibiotic therapy as prescribed to reduce the risk of antibiotic resistance. Selection of antibiotics should generally be made after wound swabs and sensitivity testing to determine the bacteria against which treatment should be directed\(^{(10)}\).

Clinical indications that the patient has spreading infection (e.g. cellulitis) include\(^{(10)}\):

- signs of advanced local infection,
- spreading erythema,
- fever,
- oedema of regional tissues, and
- general malaise

Clinical indications that the patient has systemic infection (e.g. bacteraemia, sepsis) include\(^{(10)}\):

- high fever,
- hypothermia,
- lymphangitis and regional lymphadenopathy,
- delirium,
- multiple compromise or organ failure, and
- circulatory shock (hypotension, tachypnoea, tachycardia).

All PIs should be assessed regularly for indicators of infection. For complex, unresponsive, recalcitrant or recurrent infection, consider consulting a microbiologist or infectious disease specialist\(^{(10)}\).

**Select a wound dressing.** Wound dressings or devices are applied to a wound in order to protect the wound from contamination and trauma, to absorb exudate, to fill dead space deficits, reduce oedema and to promote an optimal healing environment. Wound healing is based on the principles of moist wound healing, which is optimised through dressings that donate fluid to the wound or the application of occlusive or semi-occlusive dressings, and wound bed preparation. All products should be used in accordance with the manufacturer’s directions\(^{(10)}\).
Characteristics that are likely to influence wound dressing selection may include:\(^{10}\):

- Condition of surrounding skin,
- Ease of application and removal,
- Ability to maintain moisture balance,
- Ability to absorb exudate and odour,
- Pain experienced on dressing changes,
- Infection control and ability to maintain bacterial balance,
- Cosmetic effect,
- Skill and knowledge of the health professional,
- Accessibility and cost effectiveness,
- Suitability of dressing location to wound location, and
- Comfort.

Continually moist gauze should be used only when other moisture retentive dressings are not available\(^{10}\).

Negative pressure wound therapy (NPWT) is a wound management technique that involves application of suction to the wound using a vacuum dressing. It is reported to improve nutritional and oxygen delivery to the wound through reduction of oedema, to remove wound exudate, to promote tissue granulation, and to remove wound inhibitory factors. The therapy is primarily used to reduce wound volume and may be used to prepare the wound bed for flap closure surgery. A foam or gauze cavity filling dressing is used to fill the wound defect and sealed with a transparent film secondary dressing, through which a drainage tube connected to vacuum is inserted. When using TNPT\(^{10}\):

- Debride necrotic tissue prior to applying NPWT.
- Evaluate the wound and effectiveness of therapy with each dressing change.
- Comply with the health provider’s policies and protocols and the manufacturer’s instructions for the application, maintenance and removal of NPWT.

**A Combined Lesion**

So if there is a link between the two, do we sometimes have to treat both?

Sometimes elements of both treatments will need to be used when there is a PI present within an area of IAD. The moisture that formed the IAD may have been a contributing factor in the formation of a PI, or alternatively, the exudate from the PI may have caused the maceration/excoriation associated with the dermatitis. Either way, just managing moisture or just removing pressure will not be effective and a combined management plan will need to be implemented\(^{3}\).
## Appendix A – PI vs IAD, Comparison of Wound Related Characteristics

<table>
<thead>
<tr>
<th>Causes</th>
<th>Pressure Ulcer</th>
<th>Moisture Lesion</th>
<th>Remarks</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Pressure and/or shear must be present.</td>
<td>Moisture must be present (e.g., shining wet skin caused by urinary incontinence or diarrhea).</td>
<td>If moisture and pressure/shear are simultaneously present, the lesion could be a pressure ulcer as well as a moisture lesion (combined lesion).</td>
</tr>
</tbody>
</table>

| Location | A wound not over a bony prominence is unlikely to be a pressure ulcer. | A moisture lesion may occur over a bony prominence. However, pressure and shear should be excluded as causes and moisture should be present. A combination of moisture and friction may cause moisture lesions in skin folds. A lesion that is limited to the anal cleft only and has a linear shape is not a pressure ulcer and is likely to be a moisture lesion. Perianal redness/skin irritation is most likely to be a moisture lesion resulting from faeces. | It is possible to develop a pressure ulcer where soft tissue is compressed (e.g., by a nutrition tube, nasal oxygen tube or urinary catheter). Wounds in skin folds of bariatric patients may be caused by a combination of friction, moisture, and pressure. Bones may be more prominent where there is significant tissue loss (weight loss). |

| Shape | If the lesion is limited to one spot, it is likely to be a pressure ulcer. Circular wounds or wounds with a regular shape are most likely pressure ulcers; however, the possibility of friction injury has to be excluded. | Diffuse different superficial spots are more likely to be moisture lesions. In a kissing ulcer (copy lesion) at least one of the wounds is most likely caused by moisture (urine, faeces, transpiration, or wound exudate). | Irregular wound shapes are often present in a combined lesion (pressure ulcer and moisture lesion). Friction on the heels may also cause a circular lesion with full-thickness skin loss. The distinction between a friction lesion and a pressure ulcer should be made based on history and observation. |

<p>| Depth | Partial-thickness skin loss is present when only the top layer of the skin is damaged (Stage II). In full-thickness skin loss, all skin layers are damaged (Stage III or IV). If there is a full-thickness skin loss and the muscular layer is intact, the lesion is a Stage III pressure ulcer. If the muscular layer is not intact, the lesion should be diagnosed as a Stage IV pressure ulcer. | Moisture lesions are superficial (partial-thickness skin loss). In case where the moisture lesions get infected, the depth and extent of the lesion can be enlarged/deepened extensively. | An abrasion is caused by friction. If friction is exerted on a moisture lesion, this will result in superficial skin loss in which skin fragments are torn and jagged. |</p>
<table>
<thead>
<tr>
<th><strong>Necrosis</strong></th>
<th>A black necrotic scab on a bony prominence is a pressure ulcer (unstageable). Necrosis can also be considered present at the heel when the skin is intact and the black/blue shimmer is visible under the skin (deep tissue injury-the lesion will most likely resolve into a necrotic eschar)</th>
<th>There is no necrosis in a moisture lesion.</th>
<th>Necrosis starts without a sharp edge but evolved into sharp edges. Necrosis softens up and changed colour (eg, blue, brown, yellow, or grey) but is never superficial. Distinction should be made between a black necrotic scab and a dried blood blister.</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Edges</strong></td>
<td>If the edges are distinct, the lesion is most likely a pressure ulcer. Wound with raised and thickened edges are old wounds.</td>
<td>Moisture lesions often have diffuse or irregular edges.</td>
<td>Jagged edges are seen in moisture lesions that have been exposed to friction.</td>
</tr>
<tr>
<td><strong>Colour</strong></td>
<td><em>Red skin:</em> If redness is non-blanchable, this is most likely a pressure ulcer Stage I. For people with darkly pigmented skin, persistent redness may manifest as blue or purple. <em>Red in wound bed:</em> If there is red tissue in the wound bed, the wound is either a Stage II, a Stage III or a Stage IV pressure ulcer with granulation tissue in the wound bed. <em>Yellow in wound bed:</em> Softened necrosis is yellow and not superficial. Slough is a creamy, thin and superficial layer. <em>Black in the wound bed:</em> Black necrotic tissue in the wound bed indicates a pressure ulcer.</td>
<td><em>Red skin:</em> If the redness is not uniformly distributed, the lesion is likely to be a moisture lesion (exclude pressure and shear as causes). <em>Pink or white surrounding skin:</em> Maceration resulting from moisture.</td>
<td><em>Red skin:</em> If the skin (or lesion) is red and dry or red with a white sheen, it could be a fungal infection or intertrigo. This is often observed in the anal cleft. <em>Green in wound bed:</em> Infection. Be aware that zinc oxide ointments may result in whitened skin. While eosin is not recommended, it is still used in some areas. It will turn the skin red/brown and obstruct the observation of the skin.</td>
</tr>
</tbody>
</table>

Modified from Pressure Ulcer Advisory Panel’s “Differentiation Between Pressure Ulcers and Moisture Lesions”[1]
Appendix B – Pan Pacific Guidelines Flowchart for Prevention and Management of Pressure Injury

FLOW CHART FOR PREVENTION AND MANAGEMENT OF PRESSURE INJURY
The Pan Pacific Clinical Practice Guideline for the Prevention and Management of Pressure Injury

Assess all patients within eight hours of admission to service
Consult the patient and multidisciplinary team for care planning

CONDUCT PI RISK ASSESSMENT

Does the patient have an existing pressure injury?

YES

NO

IMPLEMENT PREVENTION PLAN

Strategies for patients at high risk/with a PI
• Implement skin protection strategies (p 37)
• Provide high protein oral nutritional supplements (Grade B, p 62)
• Consider arginine supplements (Grade C, p 64)
• Consider alternative pressure redistribution support surfaces (Grade A, p 65)
• Consider more frequent repositioning (Grade A, p 66)
• Patient education (p 55)

Preventative strategies for patients at risk
• Implement skin protection strategies (p 37)
• Consider high protein oral nutritional supplements (Grade B, p 37)
• Use constant low pressure redistribution support surfaces (Grade A, p 38)
• Regular repositioning (Grade A, p 48)
• Patient education (p 55)

ASSESS EXISTING PI

PI assessment
Use a validated pressure healing assessment scale (Grade C, p 51)

PI classification
Use NPUAP/EPUAP pressure injury classification system (p 53)

Pain assessment
Use a validated pain assessment tool (Grade C, p 57)

TREAT EXISTING PI

Pain management
• Develop an individualised pain management plan including regular analgesia (p 59)
• Consider topical opioids when debriding (Grade C, p 60)

Additional management options
• Consider electrotherapy (Grade B, p 78)
• Consider using a medical grade sheepskin as an adjunct (Grade C, p 45)

Pressure injury wound management
• Debride the wound as indicated (p 67)
• Treat infection - Consider using cadexomer iodine (Grade C, p 69)
• Select a wound dressing (p 75) - Consider negative pressure wound therapy (Grade C, p 77)

TREAT EXISTING PI

TREAT EXISTING PI

Ongoing PI risk assessment
At least weekly pressure injury healing assessment

DOCUMENT
All assessments
All management plans
All interventions

Ongoing risk assessment

APRIL 5, 2016

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### Appendix C – Debridement

<table>
<thead>
<tr>
<th>Type</th>
<th>Mechanisms of action</th>
<th>Advantages</th>
<th>Disadvantages</th>
<th>Who/where</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Autolytic</strong></td>
<td>Uses the body’s own enzymes and moisture to rehydrate, soften and liquefy hard eschar and slough using occlusive or semi-occlusive dressings and/or antimicrobial products to create a balanced moist wound environment either by donating or absorbing moisture</td>
<td>Can be used for pre-debridement, when there is a small amount of non-viable tissue. Also suitable for wounds where other forms of debridement are inappropriate. Can be used for maintenance debridement</td>
<td>The process is slow, increasing potential for infection and maceration</td>
<td>Can be done by both generalist and specialist</td>
</tr>
<tr>
<td><strong>Enzymatic</strong></td>
<td>Uses enzymes to break down a number of inorganic and organic compounds to remove eschar and slough. Requires a secondary dressing. Both Honey and Flamminal have antimicrobial properties.</td>
<td>Suitable for wounds where other forms of debridement are inappropriate. Can be used for maintenance debridement</td>
<td>High levels of exudate can damage periwound skin. Steps must be taken to prevent periwound maceration and excoriation.</td>
<td>Can be done by both generalist and specialist</td>
</tr>
<tr>
<td><strong>Mechanical</strong></td>
<td>Traditional method involves using wet to dry gauze that dries and adheres to the top layer of the wound bed, which is ‘pulled’ away when the dressing is removed</td>
<td>Newer methods are more selective, faster and relatively pain-free.</td>
<td>Non-selective and traditional methods are potentially harmful Requires frequent dressing changes and can be very painful for the patient</td>
<td>Can be done by both generalist and specialist</td>
</tr>
<tr>
<td><strong>Biosurgical</strong></td>
<td>Larvae of the green bottle fly are used to remove necrotic and devitalised tissue from the wound. Larvae are also able to ingest pathogenic organisms in the wound.</td>
<td>Highly selective and rapid</td>
<td>Costs are higher than autolytic debridement, but treatment is short once in place Not suitable for all patients or wounds</td>
<td>Can be applied by generalist or specialist practitioner with training. Closed bag method reduces skill level required and can be left for 4-5 days</td>
</tr>
<tr>
<td><strong>Hydrosurgical</strong></td>
<td>Removal of dead tissue using a high energy saline beam as a cutting implement</td>
<td>Short treatment time and selective. Capable of removing most if not all devitalised tissue from the wound bed</td>
<td>Requires specialist equipment. There is potential for aerosol spread and it is associated with higher costs</td>
<td>Must be carried out by a specialist practitioner with relevant training. Can be used in a variety of settings</td>
</tr>
<tr>
<td>Type</td>
<td>Mechanisms of action</td>
<td>Advantages</td>
<td>Disadvantages</td>
<td>Who/where</td>
</tr>
<tr>
<td>----------</td>
<td>---------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------</td>
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</tr>
<tr>
<td><strong>Sharp</strong></td>
<td>Removal of dead or devitalised tissue using a scalpel, scissors and/or forceps to just above the viable tissue level. This does not result in total debridement of all non-viable tissue and can be undertaken in conjunction with other therapies (eg autolysis)</td>
<td>Selective and quick. No analgesia is required normally</td>
<td>Clinicians need to be able to distinguish tissue types and understand anatomy as the procedure carries the risk of damage to blood vessels, nerves and tendons</td>
<td>Can be done at the patient's bedside or in clinic by a skilled practitioner with specialist training</td>
</tr>
<tr>
<td><strong>Surgical</strong></td>
<td>Excision or wider resection of non-viable tissue, including the removal of healthy tissue from the wound margins, until a healthy bleeding wound bed is achieved</td>
<td>Selective and is best used on large areas where rapid removal is required</td>
<td>It can be painful for the patient and anaesthetic is normally required. It can be associated with higher costs</td>
<td>Must be performed in the operating theatre by a surgeon, podiatrist or specialist nurses following training</td>
</tr>
<tr>
<td><strong>Ultrasonic</strong></td>
<td>Devices deliver ultrasound either in direct contact with the wound bed or via an atomised solution (mist). Most devices include a built-in irrigation system and are supplied with a variety of probes for different wound types</td>
<td>Immediate and selective. It can be used for excisional debridement and/or maintenance debridement over several sessions</td>
<td>Availability issues due to higher costs and requirement for specialist equipment. Requires longer set up and clean up time (involving sterilisation of hand pieces) than sharp debridement.</td>
<td>Must be carried out by competent practitioner with specialist training in a variety of settings</td>
</tr>
</tbody>
</table>

This table has been copied directly from “Debridement made easy”\(^{(1b)}\).
### Appendix D - Antimicrobials

<table>
<thead>
<tr>
<th>Product</th>
<th>Method of Action</th>
<th>Effect on wound healing</th>
<th>Indication/contraindication</th>
<th>references</th>
</tr>
</thead>
</table>
| Cadexomer Iodine (Iodosorb - microspherical beads of hydrophilic biodegradable starch and 0.9% (w/v) iodine) | • Highly absorbent polymer, when hydrated with wound exudate swells, releasing iodine from the cadexomer-iodine complex.  
• Iodine binds to thiol and sulfydryl groups denaturing proteins and inactivating enzymes affecting cell walls, membranes, and cytoplasmic components  
• Able to bind to fatty acids and to nucleic acids.  
• Broad microbicidal benefits | • Evidence of positive effect in burns, chronic leg ulcers, and decubitus ulcers  
• Has been shown to stimulate vascular endothelial growth factors  
• Effective in reducing bioburden in biofilms  
• Preferable to povidone iodine  
• Economical | • Contraindicated in people with sensitivity to iodides  
• highly toxic if ingested  
• Avoid contact with eyes and mucous membranes  
• May cause contact dermatitis  
• Stains  
• Resistance is rare - first documented in iodinated swimming pools and also reported in strains of Staphylococcus aureus | (16-21) |
| Silver - Nanocrystalline silver such as Acticoat or as Ionic Silver combined with many dressing types including, but not limited to, hydrofibres (Aquacel) and foams (Mepilex Ag) | • Hinders bacterial respiration  
• Disrupting the cell membrane  
• Denatures proteins  
• Alters nucleic acids  
• Effective against aerobic, anaerobic, Gram-negative and Gram-positive bacteria, yeast, filamentous fungi and viruses  
• Rapidly bactericidal | • Anti-inflammatory properties  
• Silver is good at preventing biofilm formation but not breaking down established biofilms  
• Combined with many dressing types allows for other important factors such as exudate management and odour control | • Can stain  
• Resistance to silver is rare but has been documented in chronic wounds | (22-24) |
| Honey (Manuka Honey, Medihoney) | • High osmolarity and ability to minimize water availability to bacteria  
• Slow and sustained production of hydrogen peroxide  
• Favonoids and aromatic acids demonstrate antimicrobial properties | • Effective wound deodorizing agent  
• High glucose content and low pH stimulate macrophages  
• Promotes autolytic debridement  
• Anti-inflammatory properties  
• Regulates wound pH | • Moderate efficacy against multiple drug resistant bacteria  
• Safe for use on the face and on children  
• Contraindicated in people with bee sensitivities  
• Does not have confirmed efficacy when | (19, 25-31) |
<table>
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</table>
| Hypertonic Saline - Curity: 20% solution of NaCl on a gauze dressing, can be sheet or ribbon (previously Curasalt) | • Kills by osmosis  
• Listed by the manufacturer as a wet-to-dry dressing for mechanical cleaning of wound beds | • Reduced time to healing compared to standard treatment  
• May remove new granulation tissue upon removal of the dressing  
• Mechanical debrider for removal of slough  
• Ribbon is good for packing sinuses | • Used in conjunction with compression therapy  
• Very inexpensive  
• Contains cotton products that can shed fibres into the wound  
• May cause pain on removal  
• May irritate sensitive skin  
• Can desiccate the wound surface | (32) |
| Chlorhex impregnated gauze - Bactigras | • Chlorhexidine is a biguanide, which in low concentrations inhibits enzymes associated with bacterial membranes and causes leakage of cellular material.  
• Inhibits growth of bacteria | • The efficacy of chlorhexidine against biofilms is inconclusive  
• May remove new granulation tissue upon removal of the dressing  
• Allows exudate to pass through to secondary dressing | • Non-stick  
• Contains cotton products that can shed fibres into the wound  
• May irritate sensitive skin  
• May cause pain on removal  
• Some bacterial resistance reported | (23, 33, 34) |
| Prontosan Solution (Polyhexamethylene biguanide (PHMB) (0.1%) and Betaine a surfactant (0.1%)) | • Has demonstrated efficacy in the ability to penetrate difficult to remove coatings, lifting debris, bacteria and biofilm on the wound.  
• It is thought to break down the lipopolysaccharide layer (LPS) of the bacertial wall to kill the bacteria.  
• Betaine interferes with the production of homoserine lactone a signalling molecule used in cell to cell communication of biofilms (known as quorum sensing) which play a role in biofilm pathogenicity. | Non cytotoxic, can provide an effective alternative to antibiotics and antimicrobial dressings as a method of controlling bacterial proliferation in wounds, it provides an effective method of supporting wound bed preparation and removal of biofilm | • The quick action to kill bacteria reports that PHMB is unlikely to develop resistance  
• Prontosan should not be used on anyone with a known or suspected allergy to betaine or PHMB. It is contraindicated on the central nervous sytem or meninges, middle or inner ear, eyes or on hyaline cartilage during surgery.  
• Do not combine with other wound cleansers or ointments | (24, 36) |
References


