

DECREASING NURSING WORKLOAD: SIMPLIFYING WOUND MANAGEMENT WITH AN INNOVATIVE TRANSFORMING POWDER DRESSING



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SOC TREATMENT MIX

INTRODUCTION

The current nursing shortage is expected to intensify, especially as an aging population burdens our healthcare systems¹. Limited staffing can:

- Cause safety and care issues for patients
- Negatively impact wound healing, and
- Increase hospital length of stay.^{2,3}

We investigated if using an extended-wear transforming powder dressing (TPD) would simplify wound management and decrease nursing workload by reducing requisite dressing changes and time associated with wound care, without compromising outcomes.

METHODOLOGY AND MATERIALS

Global data from 76 patients in six patient cohorts treated with TPD was aggregated. Dressing change frequency and nursing time spent on wound care using TPD was compared to standard of care (SOC) dressings. SOC dressing, dressing change time and number of weekly treatments were recorded where available or estimated based on the most conservative of three expert opinions.

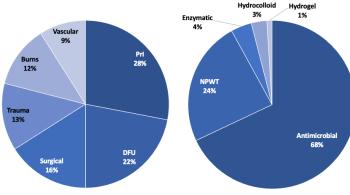
TPD is a commercially available dressing comprised primarily of hydrogel polymers like those used in contact lenses. When hydrated with saline, TPD aggregates to form a moist, oxygen-permeable barrier that covers and protects the wound while releasing excess exudate through vapor transpiration. TPD may be left in on the wound for up to 30 days and topped off as needed without requiring primary dressing changes. Simple secondary dressings may be used in areas of high exudation or friction. TPD flakes off as the wound heals.

DEMOGRAPHICS

> N: 76 | Age: 4 – 95 years

Gender: 33% female / 67% male | Wounds: 41% acute / 59% chronic

WOUND ETIOLOGIES DISTRIBUTION



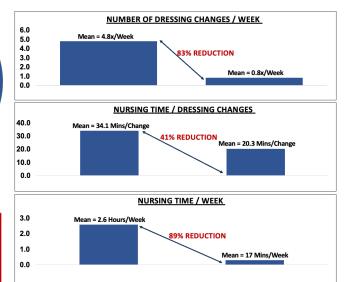
NURSING EFFICIENCIES ACHIEVED ACROSS ETIOLOGIES

WOUND TYPE	WEEKLY DRESSING CHANGE FREQUENCY			TIME / CHANGE (MINS)			TIME / WEEK (MINS)		
	SOC	TPD	% CHANGE	SOC	TPD	% CHANGE	SOC	TPD	% CHANGE
Surgical	5.6x	1.1x	-81%	34	20	-53%	154	17	-90%
Trauma	4.1x	1.1x	-73%	49	23	-53%	174	25	-86%
Burns	7.0x	0.7x	-90%	29	19.4	-33%	202	13	-94%
Prl	5.4x	0.7x	-88%	23	16	-30%	122	11	-91%
DFU	3.6x	0.8x	-77%	33	23	-31%	111	19	-83%
Vascular	2.9x	0.9x	-70%	20	14	-29%	52	13	-75%
All	4.8x	0.8x	-83%	34	20	-41%	154	17	-89%
Acute	5.5x	1.0x	-83%	46	23	-49%	223	22	-90%
Chronic	4.3x	0.8x	-83%	26	18	-30%	107	14	-87%

RESULTS

- SOC: mean dressing change frequency of 4.8x /week (3x/day to 1x/week) requiring 34 mins / change (10 -120 mins)
- TPD: mean dressing change frequency of 0.8x / week (0.2x to 2.0x/week) requiring 20 mins / change (10 -60 mins)

MEAN TIME / WEEK FOR WOUND CARE = 2.6 HOURS / WEEK WITH SOC VERSUS 17 MINS / WEEK WITH TPD



CONCLUSION

Mean dressing change frequency and required nursing time was significantly lower with TPD versus SOC for all wound etiologies. All wounds healed without complications. Pain and pain medications reduced in cases with patient reported pain. Wound management may be simplified with TPD without compromising healing outcomes by reducing dressing changes and overall costs by decreasing utilization of nursing time and material resources.

REFERENCES & ACKNOWLEDGEMENTS

(1) AACN Fact Sheet-Nursing Shortage aacnnursing.org/news-information/fact-sheets/nursing-short/ Accessed online 29OCT2022 | (2) Nursing Shortages: Implications on Wound Care and The Way Forward. April 5, 2022. Healiant Training & Education. https://healiant.com/nursing-shortages-implications-on-wound-care-and-the-way-forward | (3) Nurse staffing models, nursing hours, and patient safety outcomes. nlm.nih.gov. age. Accessed online 28OCT2022.

Acknowledgements: This poster was created in collaboration with Altrazeal Life Sciences Inc. (ALSI). Mr. Ron Sototmayor and Dr. Johnathan Saxe serve as clincial consultants to ALSI. Dr. Saxe also served as ALSI's Medical Director. | EDU-0076, Rev 01

trazeal® transforming powder dressing Simplify Wound Healing - Skin 2.0

TRANSFORMING POWDER DRESSING WITH EXTENDED WEAR TIME **RETROSPECTIVE EVALUATION IN TRAUMA INJURIES**

Jonathan M Saxe, MD, MAR, MBA, FACS; Vice Chair of Surgery; Ascension St. Vincent Medical Center, Indianapolis, IN; Medical Director, ULURU Inc.

MTEC Annual Membership Meeting, Baltimore, MD | May 2022

COMPANY MISSION | ACHIEVEMENTS TO DATE

OVERVIEW: Based in Addison, TX, we are committed to developing and commercializing innovative wound care and drug delivery systems based on our patented technologies.

MISSION: To improve the lives of patients the world over by delivering comprehensive solutions that optimize outcomes for all key stakeholders: patients, providers and payers.

ALTRAZEAL STATUS: Patent granted, FDA registered, and market launch initiated. Used in prominent hospital systems: AdventHealth, Northwestern, Northwell Health, NYU, Plaza Medical (HCA), Providence Health (LifePoint), St. Vincent's (Ascension), UPMC.

DOD / VA EFFORTS: DAPA listed. SAM registered. Introduced to MRMC. CCCRP. USAISR. USUHS. WRAIR, NMRC, 59th Wing. Approved for use in several VAMCs including Boston, Chicago, Charleston, Dallas, Sacramento and Tampa,

DOD FUNDED R&D PROJECTS: Received grant funding from the Department of Defense related to three post-marketing clinical studies as well as a number of pre-clinical studies for product development:

1. MTEC-NAMD: Clinical studies in diabetic foot ulcers and acute partial thickness burns

2. CDMRP-DHA: Clinical study in pressure injuries

3. SBIR-WRAIR-DHA and MIDRP-BDRD: Development of Altrazeal combinations for drug delivery

UNIQUE EXPERTISE: Clinical partnerships with global wound care experts and centers of excellence.

ABOUT ALTRAZEAL® TRANSFORMING POWDER DRESSING

Altrazeal powder is comprised primarily of two biocompatible polymers (same as those used in contact lenses). Upon hydration, Altrazeal granules aggregate to form a moist, oxygenpermeable barrier that protects the wound from contamination while helping to manage excess exudate through vapor transportation. Once applied, Altrazeal may be left in place for up to 30 days. Additional powder may be added ("topped off") as needed without requiring primary dressing changes. Simple secondary dressings may be used in areas of high friction or exudation. If clinically necessary, the dressing may be removed atraumatically by lifting off with a pair of forceps. As the wound heals, Altrazeal dries and flakes off.

Altrazeal[®] Pain Scores Significantly Lower at All Time Points in the Randomized Study



Dressing Changes: > Total: Two changes

- > Average: Once every five days
- Outcomes:
 - > Wound healed in 10 days with two applications
 - > Avoided grafting

CASE SERIES OVERVIEW: FULL THICKNESS TRAUMATIC INJURIES (N=10)

A retrospective analysis of data from 10 patients with complex, full thickness traumatic wounds with elements of exposed bones, tendons and muscle was performed. All wounds were treated with Altrazeal until complete wound closure was achieved or the wounds were ready for grafting. Time to healing as well as frequency and number of dressing changes were tracked.

CASE 1: TRAUMA INJURY FROM MOTORCYCLE ACCIDENT

- Patient: 31-year-old female
- Wound Size: 35cm x 17cm x 17cm
- · Challenge: Extensive wound with dressing changes required multiple times a week
- Dressing Changes:
- > Total: Six changes
- > Average: Once every eight days

· Patient: 40-year-old male

 Dressing Changes: > Total: Two changes

Outcomes:

Wound Size: 25cm x 25cm x 5cm

> Average: Once every nine days

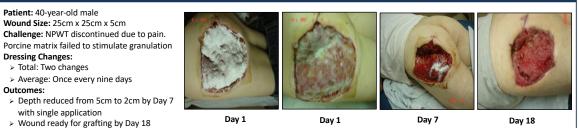
with single application

- Outcomes:
- > Wound healed in 45 days > Avoided grafting

Day 37 Day 1 Day 30

Day 4 Day 8 Day 45





CASE 3: GUN SHOT WOUND



CONCLUSION

A marked acceleration in healing was observed in all ten cases with reduced frequency of dressing changes relative to standard of care (once every 6.5 days on average versus daily or three times a week for standard of care). Six patients achieved complete healing without requiring grafting. Mean healing time was 23 days with three applications on average, or once every seven days. Four patients received successful grafting treatments once sufficient granulation had been achieved with Altrazeal. Average treatment time was 33 days with dressing changes every six days on average. Clinical observations and outcomes indicate that Altrazeal presents a safe, effective and resource-efficient modality for the treatment of patients with traumatic wounds.



A Novel Pre-Grafting Wound Management Technique to Promote Granulation in Complex Painful Wounds



Jonathan M. Saxe, MDMAR, MBA, FACS; Ascension St. Vincent Hospital, Indianapolis, IN, USA; Mounir Mabrouk, MD; Alexandria University, Egypt | WOCNext 2022 Meeting, Fort Worth, TX, June 5-8, 2022

BACKGROUND

Skin grafting (SG) is used to provide coverage in both acute and chronic wound settings. Preparation of the wound bed with development of granulation tissue is vital for graft success.¹ Traditional standard of care (SOC) wound management principles involve debriding the wound followed by negative pressure wound therapy (NPWT), bolstering or conventional dressing applications to accelerate wound healing prior to grafting.² Current SOC is limited in complex painful wounds. NPWT deployment and application is often difficult and painful.³ Pain is also a significant issue often associated with repeated wound dressing changes.

METHODOLOGY & MATERIAL

Three case studies incorporating treatment with Transforming Powder Dressing (TPD) to promote granulation in patients with complex wounds are reviewed. In each case, patients had extensive wounds with high levels of reported pain. Prior treatment in all cases failed to progress wounds to the stage to permit grafting. Cases reviewed include a large abdominal wound resultant of an automobile accident, complex wounds associated with uncontrolled chronic vasculitis, and an improvised explosive device (IED) accident. TPD was introduced to facilitate and promote granulation and allow grafting.

TPD is comprised primarily of biocompatible polymers. Upon hydration with saline, TPD granules aggregate to form a moist, oxygen-permeable matrix that protects the wound from contamination while helping to manage excess exudate through vapor transpiration. Once applied, TPD may be left in place for up to 30 days and additional powder may be added ("topped off") as needed without requiring primary dressing changes. Simple secondary dressings may be used in areas of high exudation or friction. TPD dries and flakes off as the wound heals.

RESULTS

PATIENT 1: 40 y/o female with large complex abdominal wound resultant of car accident

Challenge: NPWT could not be placed due to risk of fistula **TPD Treatment:** TPD applied to wound with foam for excess exudate absorption

- **Outcomes Post-TPD Treatment:**
- Wound was ready for grafting by Day 18

PATIENT 2: 42 y/o male with non-healing progressive venous ulcer, uncontrolled vasculitis, infection history, necrosis, exposed bone

Challenge: High pain score (9/10), failed SOC treatment Treatment: TPD 2x/week in the first week and 1x/week after Outcomes Post-TPD Treatment:

- Significant reduction in pain
- Granulation tissue covered bone and patient was grafted
 Avoided amputation

PATIENT 3: 40-year-old male with a 25cm x 25cm x 5cm IED blast wound due to consumer firework accident (M-80) Challenge: NPWT was discontinued due to patient pain, porcine matrix failed to stimulate granulation Treatment: TPD applied weekly

Outcomes Post-TPD Treatment:

Wound depth reduced from 5cm to 2cm by day 7
Wound was ready for grafting by day 18

CONCLUSION

A marked acceleration in granulation was observed in all three cases. Patients reported reduced pain and the frequency of dressing changes were also reduced relative to SOC. No adverse events were reported. Based on the clinical observations and outcomes, we conclude that TPD presents a safe and effective modality for preparing complex painful wounds for successful grafting.







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Novel Technique for Management of Painful Road Rash Injuries

Lori O'Shea, BSN, RN, WCC; Jenny A. Ziembicki, MD, FACS | UPMC Mercy Hospital Burn Center

Symposium on Advanced Wound Care (SAWC) | April 2022



BACKGROUND

Road rash injuries include painful skin abrasions, burns or wounds resulting from trauma accidents on cemented or tarred surfaces. Wounds vary in severity, depth, and degree^{1, 2}

- Complications include infection, sepsis, wound progression, pain (often significant, resulting from wounds and wound related treatments^{3,4}), embedded road debris, and devitalized tissue.
- Routine Standard of Care includes topical antibiotics, petroleum dressings, or moisture retaining therapies, and requires frequent and painful dressing changes.

CASE OVERVIEW: METHODOLOGY

20-year-old male sustained multiple injuries in a motorcycle accident, including:

- Subarachnoid head bleed
- Multiple mixed partial deep and superficial thickness wounds on his arm and legs

Initial treatment:

 Primarily focused on patient head injury and maintenance of neurological stability, which precluded use of pain medications. Because cleaning of wounds was reported as highly painful, all wounds were initially managed conservatively with simple petroleum-based contact layers and absorbent pads.

Upon hospital discharge 4 days later:

- He was treated at home by his caregiver, a wound care nurse. Due to the high level of pain, his caregiver elected to utilize TPD* to treat the wounds instead of application of topical antibiotics. Mechanical and autolytic debridement was performed to manage the exudate and remove the remaining embedded road debris.
- TPD was applied (sprinkled on the wounds), followed by a contact layer and gauze after the wounds were cleaned.

MATERIALS

TPD is a novel powder dressing comprised primarily of biocompatible polymers. Upon hydration with saline, TPD granules aggregate to form a moist, oxygen-permeable matrix that protects the wound from contamination while helping to manage excess exudate through vapor transpiration, as well as some negative pressure effects on the wound. Once applied, TPD may be left in place for up to 30 days and additional powder may be added as needed without requiring primary dressing changes. Simple secondary dressings may be used in areas of high exudation or friction. TPD dries and flakes off as the wound heals.

TPD APPLICATION AND RESULTS

- TPD was topped off as needed, secured with contact layers, absorbent pads and a net dressing.
- On post-injury day nine, five days after initial TPD application, all wounds were epithelialized.
- Immediate pain relief was reported by the patient following TPD application.
- There were no complications of infection or wound progression.
- · Post-healing scarring was minimal to absent.



CONCLUSION

Based on the accelerated wound healing, and pain reduction results associated with this patient, we conclude that TPD offers a safe and effective alternative to Standard of Care for the management of painful road rash injuries.

REFERENCES AND ACKNOWLEDGEMENTS

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The Utility of Transforming Powder Dressing in the Treatment of Various Wound Types: A Case Series



Sawyer Cimaroli, MD; Avi Hatami, MD; Brian Gillette, PhD; Scott Gorenstein, MD | NYU Langone Long Island Hospital, Department of General Surgery SAWC Fall 2022 Meeting, Las Vegas, NV | October 13-16

INTRODUCTION:

More than 8 million people a year receive wound treatment in the U.S. Increasing health care costs, limited healthcare resources, an aging population, and life-style related diseases make wound management a growing clinical, social, and economic burden, indicating a vital need for a more effective wound management solution.

Transforming powder dressing (TPD*) forms a non-occlusive barrier which helps maintain a moist environment while facilitating the flow of excess exudate via vapor transpiration.



METHODOLOGY & MATERIALS:

- TPD is a novel, biocompatible polymer powder dressing which transforms into an extended-wear gel matrix upon hydration
- Single-center retrospective case series of various acute and chronic wounds treated with TPD
- 50 patients with 50 wounds were treated with TPD and had at least one follow-up visit during the study period

SUMMARY OF DEMOGRAPHICS				
Number of subjects	50			
Number of TPD applications	129			
Mean age (range)	76 years (18-99)			
Male to female participants	20 (40%) / 30 (60%)			
Number of current smokers	3 (6%)			
Number of previous smokers	25 (50%)			
Mean number of comorbidities (range)	6 (0-15)			

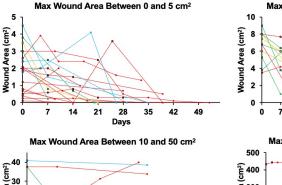
*Altrazeal[®] Transforming Powder Dressing

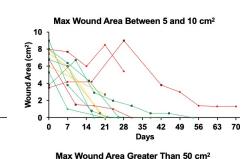
TREATMENT:

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SUMMARY OF WOUND HEALING						
WOUND TYPE	TOTAL NUMBER	MEAN NUMBER OF TPD APPLICATIONS	NUMBER WITH WOUND HEALING > 10% PER WEEK**	NUMBER HEALED		
Venous Leg Ulcer	27 (54%)	2.8	17 (63%)	12 (44%)		
Traumatic Wound	11 (22%)	2.5	10 (91%)	6 (55%)		
Skin Tear	7 (14%)	2.6	5 (71%)	5 (71%)		
Other (Surgical, Arterial, Diabetic)	5 (10%)	1.6	4 (80%)	4 (80%)		
TOTAL COUNT / MEAN	50	2.6	36 (72)%	27 (54%)		

** Based on average healing per week from baseline to last visit





120 150 180 210 240 270 300

Davs

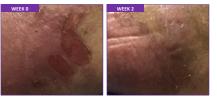
Other • Visit • Altrazeal Application

ILLUSTRATIVE CASES:

Case 1: Venous Ulcer



Case 2: Trauma | Skin Care



DISCUSSION:

Wound area reduction and positive healing trajectories were observed in nearly all wounds regardless of etiology. Complete healing was achieved in 54% of all wounds with 2.6 mean TPD applications. No TPD related adverse events were recorded.

Chronic Wounds: Complete healing was achieved in 44% of VLU treated with TPD and compression with 2.8 mean applications. TPD was ideal for venous leg ulcers because of its absorptive property and ability to conform and adhere to challenging locations.

Acute Wounds: 61% traumatic and skin tear wounds healed completely with 2.5 mean applications.

We conclude that TPD is a safe and effective way to treat acute and chronic wounds of various etiologies and locations.

REFERENCES & ACKNOWLEDGEMENTS:

21

Davs

28

VLU — Traumatic Wound

35

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Skin Tear

g 300

D 200

30 60 90

ຊັ້ 100

Acknowledgments: The study was conducted independently by NYU and no compensation was paid to the authors. This poster was presented in collaboration with ULURU Inc. For application instructions and risks of this device refer to Altrazeal Instructions for Use.

EDU-0048, REV 01



Effective Treatment of Anticoagulated Patients with Novel Transforming Powder Dressing (TPD)

Tammy Lichtman, RN, BSN, CWON; Jeffrey Chiu, MD; Ron Sotomayor, BA, RN, CWOCN; AdventHealth System, Orlando, FL | Symposium on Advanced Wound Care (SAWC) Spring Meeting, April 2022

Background

Anticoagulants can contribute to wound healing complications by accelerating bleeding and exudation. Even minor injuries may evolve into severe soft tissue damage with increased risk of hematomas. Resulting wounds may be extensive, painful, and debilitating with high risk of infection and tissue necrosis, imposing a significant wound care challenge with limited treatment alternatives.¹ Conventional dressings require frequent changes exacerbating wound trauma and patient discomfort. Another routine therapy, negative pressure wound therapy (NPWT), has been associated with bleeding complications. Between 2009 and 2011, the FDA reported six deaths and 77 injuries associated with NPWT; bleeding complications were prevalent in all six deaths and 17 injuries, including certain patients on anticoagulant therapy.² Alternative therapies, therefore, must be evaluated.

Case Overview: Methodology

We present a complex case involving a patient treated with a novel transforming powder dressing (TPD*). An 88-year-old male with multiple comorbidities including atrial fibrillation was being treated with apixaban to reduce the risk of stroke and systemic embolism. The patient reported a fall, wounding his right lower extremity (RLE). His pain increased and the wound worsened despite standard of care therapy. Computed Tomography Angiography revealed a large, superficial hematoma in the lateral aspect of the right calf. Acute blood loss and anemia secondary to the hematoma was observed. DVT prophylaxis was discontinued. Excisional wound debridement and evacuation of the hematoma was performed eleven days after the initial injury. TPD was applied to the wound post-surgery and continued during home healthcare visits. The patient was observed weekly for removal of contact layer and outer dressing to assess the wound. For the first 5 weeks, the wound dressing was changed 1 time every other week. The final 6 weeks, the wound dressing was changed 1 time every third week (11 TPD dressing changes over treatment course).

Material

TPD is a novel powder dressing comprised primarily of biocompatible polymers (same as those used in contact lenses). Upon hydration with saline, TPD granules aggregate to form a moist, oxygen-permeable matrix that protects the wound from contamination while helping to manage excess exudate through vapor transpiration, as well as some negative pressure effects on the wound. Once applied, TPD may be left in place for up to 30 days and additional powder may be added ("topped off") as needed. Simple secondary dressings are used and changed in areas with high exudation or friction. TPD dries and flakes off as the wound heals.

*Altrazeal® Transforming Powder Dressing (USA)

Results

- After treatment with TPD our team observed the following results:
- Reduced pain, minimal wound bleeding and a significant decrease in sanguineous drainage
- Expedited granulation: wound was ready for grafting 35 days after initial application of TPD (clinical team elected not to perform grafting)
- Significant reduction in wound size and dressing changes
- From 155 cm3 to 1.2 cm3 within 118 days
- Dressing change frequency reduced relative to standard of care dressings
- Changes every 10.7 days on average
- Homecare nursing visits reduced to once per week instead of thrice per week (standard of care)



Conclusion

After treatment with TPD, we observed a significant reduction in sanguineous drainage and overall wound size. Dressing change frequency and home health visits were reduced relative to standard of care therapies.

The wound care team recommended secondary intention wound healing with TPD as grafting was no longer required.

Based on the results of this complex case, we conclude that patients who are at high risk for stroke and embolism and are being treated with anticoagulants should be considered for wound care treatment with TPD.

References

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- Livingston, M; (November 20, 2009); FDA Preliminary Public Health Notification: Serious Complications Associated with Negative Pressure Wound Therapy Systems (Shuren JE); https://woundblog.wordpress.com/2009/11/20/fda-preliminary-public-health-notification-serious-complications-associated-with-negative-pressure-wound-therapy-systems/

Acknowledgements: This poster was developed and presented in collaboration with ULURU Inc.



A Novel Transforming Powder Dressing for Healing Chronic Wounds of Multiple Wound Etiologies



David Bickers CRNP, CWOCN-AP | University of Pittsburgh Medical Center, Altoona PA WOCNext 2022 Meeting, Fort Worth, TX | June 5-8, 2022

CHALLENGE

Delayed wound healing results from an imbalance occurring during healing stages, often resulting in conversion of an acute wound to a chronic non-healing wound.^{1,2} Chronic wounds are significantly more complicated to heal than acute wounds.² In the US alone, chronic wounds currently affect 6.7 million people, with annual healthcare costs exceeding 50 billion dollars.³

Evidenced based clinical principles for optimizing wound healing include: (1) maintaining a moist (but not wet) wound environment, (2) permitting gaseous and fluid exchange while providing mechanical and bacterial protection, and (3) utilizing a dressing that is non-adherent to the wound, easy to use, comfortable and pain-free for the patient. When standard of care (SOC) therapy fails to heal a wound, alternate treatment strategies must be considered.

METHOD AND MATERIALS

We present a case series which evaluates the clinical outcomes of 3 patients with chronic wounds of different etiologies which were refractory to prescribed SOC therapy (burn, 2 diabetic foot ulcers and trauma wound).

All wounds had deteriorated or showed no clinical progress prior to conversion from SOC dressings to Transforming Powder Dressing (TPD). For purposes of consistency in our assessment, the conversion of the primary dressing from SOC to TPD was the only wound treatment factor modified.

TPD is a novel powder dressing comprised primarily of biocompatible polymers (same as those used in contact lenses). Upon hydration, TPD granules aggregate to form a moist, oxygenpermeable matrix that protects the wound from contamination while helping to manage excess exudate through vapor transpiration. Once applied, TPD may be left in place for up to 30 days and additional powder may be added ("topped off") as needed without requiring primary dressing changes. Simple secondary dressings may be used in areas of high exudation or friction. TPD dries and flakes off as the wound heals.

SUMMARY RESULTS

In the cases presented, each of which was refractory to SOC therapy, all wounds healed and came to complete closure after treatment with TPD.
Average time to heal for all 4 wounds after initial treatment with TPD was 47 days.

PATIENT 1: BURN

INITIAL VISIT

CONCLUSION

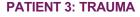
- History: 62 y/o male with DMT2, venous insufficiency, mild lymphedema, and deep partial thickness burn on ankle / LLE after catching sock on fire while welding
- Wound Size: 1.5cm x 1.5cm x 0.2cm
- Wound Duration: > 8 weeks (60 days)
- Prior Treatment: Silver Sulfadiazine 1% cream and non-adherent dressing multiple times a week
- TPD Treatment: Weekly applications
- · Outcome: Fully healed in 42 days with TPD

DAY 1: TPD APPLICATION

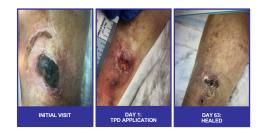
PATIENT 2: DIABETIC FOOT ULCER

- **History:** 62 y/o female with IDDM T2, lymphedema,
- neuropathy, BMI 45.6, and two plantar DFUs
- Wound Size: 0.5cm x 0.5cm x 0.7cm (heel) | 1.6cm x 1.2cm x 1.2cm (5th metatarsal)
- Wound Duration: ~1.5 to 2 years
- Prior Treatment: Total contact cast with foam dressings
- TPD Treatment: Weekly applications
- Outcomes: Both ulcers fully healed within 35 days (average)
 - Heel Ulcer: Closed in 33 days with TPD
 - Submetatarsal 5 Ulcer: Closed in 37 days with TPD





- History: 52 y/o male with CAD, renal disease, smoking disorder, and trauma wound to anterior knee
- Wound Size: 2.5cm x 2cm (eschar)
- Wound Duration: > 6 weeks (45 days)
- Prior Treatments: Mupirocin calcium ointment, medical grade honey, cortisone applied multiple times a week
- TPD Treatment: Weekly applications
- Outcomes: Fully healed in 63 days



The use of TPD as a universal primary dressing on non-healing wounds of different etiologies significantly improved healing times with reduced frequency of dressing changes and brought each of the non-healing wounds to complete closure. No adverse events were reported.

REFERENCES AND ACKNOWLEDGEMENTS

DAY 42: HEALED

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PRELIMINARY EVALUATION ON THE APPLICATON OF A POWDER DRESSING IN THE FIXATION AND TREATMENT OF CHRONIC WOUNDS **RESULTING FROM TRAUMA WITH EPIDERMAL SEPARATION**

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Purpose

The purpose of this evaluation was to test a protocol in the management of chronic wounds resulting from trauma with epidermal separation combining a novel powder dressing for both fixation and moist wound healing.

Background

A common wound seen in wound care centers is a chronic wound resulting from trauma with epidermal skin separation. This primary acute wound is commonly called a skin tear and is typically found on elderly patients. This wound often becomes chronic if not treated properly. Skin tears have been identified as a common, acute injury sustained by the aged in extended care facilities ¹⁻³ and changes to aging skin make this population more susceptible to skin tear injuries ^{1, 3-9}. A skin tear is defined as "A laceration of the epidermis, most often associated with minor trauma and involving a separation of the epidermis from the underlying connective tissue so that a flap of skin is created".

Methods:

Six patients were evaluated in a case series without controls. Patients ranged in age from 62 to 93 with a mean age of 76 years. For each patient, wounds were assessed and treatment history was documented. In all cases the trauma with epidermal separation occurred prior to referral to the wound center. The wounds were cleaned and the powder wound dressing was applied. The intact dressing was covered with a contact layer followed by a rolled gauze dressing and/or ace wraps for compression if indicated ... Patients were assessed at approximately 7 day intervals at which time the dressings were changed and the wound size and appearance were evaluated.

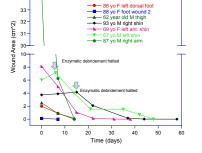


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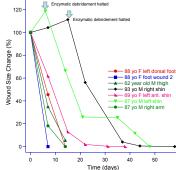


Day 28 after weeklv evaluation and dressing changes



Wound Area vs. Time

Wound % of Original Size vs. Time



Results

For the patients evaluated, six (6) patients had a total of seven (7) wounds that healed using the powder dressing. No wounds failed to heal,. The data was not generated as a prospective clinical study with inclusion criteria. Data are shown below for wound healing with and without preriods of enzymatic debridement:

Parameter	Post Enzymatic Debridement	Debridement Included
Minimum Healing Time	7 days	7 days
Maximum Healing Time	36 days	48 days
	19.8 days	24.6 days
	14 days	14 days
Std Deviation	10.5	16.4

Two patients showed an initial increase in wound area. These increases occurred with the application of an enzymatic debriding agent in conjunction with the powder wound dressing.

Of the six patients, only two reported pre-treatment pain. Both reported a decrease in pain after the application of the powder wound dressing. One reported a change from 8 to 1 with the application of the dressing on a pain scale of 1-10 (10 being the worst) and the other 5 to 0.

In all cases the dressing was applied without adhesive fixation and wrapped loosely in cotton gauze. The dressings were changed on a weekly schedule with the exception of two 14 day intervals when patients were not present for weekly dressing changes.

The dressing remained in place for all patients during the evaluation period.

Conclusions

The data shows that the powder dressing can be safely applied to a skin tear and used to anchor that tear during a treatment protocol with a 7 day interval between evaluations. A clinical study has been implemented using randomization and controls for the treatment of skin tears using this treatment protocol.

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Comparison in Management of Large, Open Combat Wounds in Service Personnel Using Negative Pressure Wound Therapy as Standard of Care and a Novel Powder Dressing

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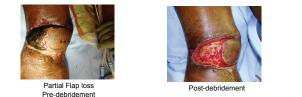


Saphenous artery fasciocutaneous flap coverage of a posterior popliteal wound with exposed nerve from a rocket-propelled grenade attack in Iraq.

Introduction

Combat trauma produces wound patterns that are seldom observed in civilian hospitals and require complicated surgery and post-operative care. The damage created by explosive devices depends on a number of factors, including the type of explosive, and the environment within which the detonation takes place. Injuries are dependent on the power of the explosion, the proximity of the casualty to the explosion and the environment (open or confined). When the energy of an explosion is directed towards tissue it creates a wound that is catastrophic to deep tissue and that will require surgical reconstruction until fully healed. In addition, this type of wound is universally accompanied by penetrating shrapnel which carries bacteria deep into the wound. Currently blast wounds in Uniformed Services Hospitals are managed uniformly with negative pressure wound therapy (NPWT), which is used to stabilize the wound, remove bacteria from tissue, and ultimately provide a wellvascularized granulation bed suitable for subsequent grafting.

This set of cases details the post-operative management of three combat trauma wounds where a combination of NPWT and a novel powder dressing are used to achieve closure. Patient is a healthy 38 year old male with a concominant closed head injury. Initial treatment was serial debridements with VAC coverage for ten days. The procedure was complicated by superficial skin necrosis which was managed by debridement and NPWT (4 days) followed by skin grafting covered with the powder dressing for 10 days. Patient is now five months post-injury with a healed wound pending secondary nerve grafting.



2 weeks post grafting

Latissimus dorsi myocutaneous free-tissue transfer with large skin grafts to the wound 2 weeks following injury. Patient is a 26 year old male healthy medic who suffered an open fracture of his anterior knee/thigh with large soft tissue loss. Initial pre-flap management was serial debridement and VAC dressings. Post-flap dressings were NPWT as a bolster over the recipient graft sites and the powder dressing to his skin-graft donor sites and the recipient site following removal of NPWT. The graft sites required two additional applications of the powder dressing until complete re-epithelialization occurred at 3 weeks following flap procedure.

Patient is now 3 months post-op with a healed wound and ambulatory. He is also currently being treated by osseous distraction/lengthening of his femur fracture. Patient will require subsequent tendon grafting to improve knee extension power.



Anterolateral thigh free-tissue transfer and large skin grafting five months ago. Patient is a 22 year old male healthy infantry soldier who suffered an IED blast injury to his lower leg and ankle with an open ankle fracture,

exposed tendons and a segmental nerve deficit. The other injury was a traumatic below-knee amputation of the opposite leg. Initial management was serial debridement, NPWTfor 10 days prior to free flap.

Post-op management was NPWT bolster over skin grafted areas with transition to powder dressing for two weeks (three applications required). Patient is completely healed, ambulatory and awaiting nerve grafting





Post Grafting

One week post free flap, powder wound dressing in place

Discussion

Complicated trauma cases involve detailed assessment and planning prior to reconstructive surgery. The choice of techniques for post-operative management is critical and can often be overlooked after the surgical procedure is completed. For trauma cases involving reconstructive surgery with flaps or skin grafts, the conventional choice is NPWT or bolster dressing(s). These techniques are viable and functional options. A novel powder dressing was evaluated as a choice for covering and protecting a flap or mesh STSG after the application of NPWT. This technique was evaluated to determine if it was possible to transition from NPWT to the use of a unique powder dressing for closure. The data set is limited, however, the powder dressing does appear to cover and protect a wound during closure with dressing changes at intervals between 3 and 7 days.

Conclusions

It is possible to transition from NPWT to the use of a novel powder dressing in post operative care of combat trauma cases involving reconstructive surgery. Further studies as to the timing of the transition would benefit the development of best clinical practice for the use of this novel dressing. Gregory A. Bohn, MD, FACS, Medical Director Trinity Wound Care and Hyperbaric Medicine Bettendorf, Iowa

Powder Dressing with high Moisture Vapor Transpiration Rate may have physical influence on wound healing

Background: Hydrogels and hydrocolloid dressings have many favorable properties and improve wound healing while maintaining a moist wound environment (4). A new powder dressing conforms to the wound surface and has a high Moisture conforms to the wound surface and has a high Moisture Vapor Transpiration Rate (1). Although the precise machanism of action has not been determined, it is transpiration nate (WTR) creates to slow pressure at the interface between the dressing and wound bed that stimulates the formation of healthy argunulation tissue (2). This novel new dressing was plexed on a post surgical for wound that, had not shown improvement with standard wound care.

Methods:

Standard wound measurements are recorded weekly as part of the patient's wound documentation. The timing of application of powder dressing technology to the wound was evaluated by observed change in wound volume over time. Wound volume was calculated using the over time. Wound volume was calculated using the formula: Wound Volume = Length x Witht x Depth x.8. This data was then displayed in graph form (Figure 1) to observe the change with the application of this new dressing product.

Results:

Nesurus: The patient's wound did not demonstrate healing as measured by wound volume calculations in the first 4 weeks of wound care. The wound volume decreased precipitously after application of this powder dressing. The wound showed a decrease in volume measurement by \$2.6% in the first week of dressing therapy. Volume culturion encode used on the theraps with reduction continued in the second week of therapy with the wound volume recorded measuring a 95.3% reduction in wound volume compared to the patients initial visit measurements. If the dressing creates a low pressure at the dressing - wound interface by way of the high Moisture Vapor Transpiration Rate promoting healthy granulation tissue formation, this dressing may stimulate healing by influencing physical characteristics of cells.

Case Study:

Case Study: 39 yo white male had suffered crush injury to the dorsum of his left foot. A split thickness skin graft was applied to cover the acute crush injury. Recent orthopedic corrective surgery was performed and resulted in a new skin and tissue deficit on the dorsum of his foot. He was referred to wound care to "clean up the wound in preparation fo a skin graft". After 4 weeks of standard wound care, the wound had stagnated as tracked using standard wound volume measurement calculations. Powder dressing became available in our clinic and was applied to the wound. The dressing remained in place and was wound. The dressing remained in place and was changed weekly at his clinic visit. Debridement was performed at each clinic visit. This patient experienced a rapid healing trajectory as demonstrated in Figure 1. His wound went on to heal without requiring an additional skin grafting procedure.

Initial Wound





Week 8





Week 5 Post-debridement

Week 8

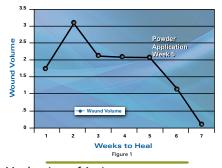


Week 5 Altrazeal

Healed



Powder Impact on Healing



Mechanism of Action – Exudate Management



Fluid Content (%)

Tensile Strength (MPa)

Conclusions: Cunctustors: Powder dressing applied to this wound promoted a vigorous granulation response and healed a wound expected to require a skin graft. The wound had not demonstrated significant progress in 4 weeks sufficient to predict healing based on well accepted parameters (7). After application of powder dressing, the wound started on a steep healing trajectory (fig. 1). This wound progressed and closed without requiring an additional operative

procedure to aim graft. Micro-stress on cell walls con cause scientiar profiferation and regrontranspiration rate of healing (3). Ghum the high moisture regrontranspiration rate of this metrical compared to other dressings, 12,000 gm/24hrs vs. 1100 gm/24hrs (2), the dressing may have a physical affect on the profiferating cells, wound fibroblasts in the wound bad and the wound margin to promote granulation formation can healing. Once the dressing is formed, the particules crease capitary forces that pull moisture that the standard standard standard standard standard standard standard freesing surface of moisture variance of all initia standards produces stress on cells and brings about conversion into rapidly metabolizing includiasts (avand frechostas). Further work to therbiblists and granulation tissue formation with subsequent wound healing would their ju understanding the affects brought about with this dressing under ju numbers and the stress of surfaces and the stress the stressing material.

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