Effects of a Novel Non-Biologic Desiccant to Remove Candida and Dermatophytes **Using a Deep Partial Thickness Wound Porcine Model**



Abstract:

Prevalence of candida and dermatophyte infections is increasing.^{1,2} Trichophyton rubrum is one of the most prominent human pathogenic dermatophytes, and accounts for almost approximately 70% of chronic dermatophytosis in humans. Candidiasis is mostly caused by Candida albicans and produce major complications in burn wounds.^{3.} Candida and dermatophyte species cannot be overlooked as an infectious entity in chronic wounds and clinicians should be focused as well as bacteria biofilms. Burn wound infections remain the most important factor limiting survival in burn intensive care units. Large wounds, impaired immune systems and broad-spectrum antibiotic therapy contribute to the growth of opportunistic fungal species.⁴ The aim of this study was examine the effects of a novel debridement method to remove necrotic tissue and dermatophytes using a porcine model.^{5,6} Deep partialthickness wounds (63) were created and inoculated with either Trichophyton rubrum ATCC28188(TR), Trichophyton interdigitale ATCC9533(TI) or Candida albicans ATCC64550(CA). Colonization was allowed by 72 hours then baseline wounds (3) were assessed prior apply treatments: 1)Desiccant Shock Technology [DST*], 2)Clotrimazole 1% Positive Control+, or 3)Gauze with sterile saline. Wounds were treated (30seconds) and then rinsed with 5ml of saline. Sterile gauze was used to remove slough. Biopsies (6mm) were taken 20minutes and 24hours for microbiology assessments. DST treated wounds showed the lowest CA64550, TI9533 and TR28188 counts at 20 minutes and 24 hours as compared to positive and untreated controls. When challenging wounds treated with CA64550, both DST and Clotrimazole 1% exhibited large fungal differences after 24 hours at 99.85% and 99.78%, respectively. Only those wounds treated with DST showed a large fungal difference when compared to baseline wounds, having 96.22% and 94.88% reductions against TI9533 or TR28188, respectively. At 24 hours when comparing DST to untreated control there was a 2.5 LogCFU/g, 2.3 LogCFU/g and 3.5 Log CFU/g reduction with CA, TI and TR, respectively. Overall, wounds treated with DST showed lower fungal counts against the three microorganisms at both 20minutes and 24hours. Comparing the Clotrimazole 1% to untreated, we observed a good reduction in fungal counts when treating wounds infected with Candida albicans ATCC64550. However, Clotrimazole 1% did not appear as effective in wounds infected with either TI9533 or TR28188. Additional animals are currently planned to substantiate these findings which may have important clinical implications in the acute and chronic wound care therapies.

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Introduction:

Infections with *C. albicans* and dermatophytes have been increased every years. ² The presence of those organism in wounded area had implications for patients. ³ Debridement techniques have shown limited ability to mechanically remove bacteria from a wound bed.¹ DST* is a topical formulation that can be used by healthcare practitioners for wound cleansing. The purpose of this study was to evaluate the ability of DST* to remove fungal infections in wounds inoculated with *Trichophyton rubrum* ATCC28188(TR), *Trichophyton interdigitale* ATCC9533(TI) or Candida albicans ATCC64550(CA).

Materials and Methods: **1. Experimental Animals:**

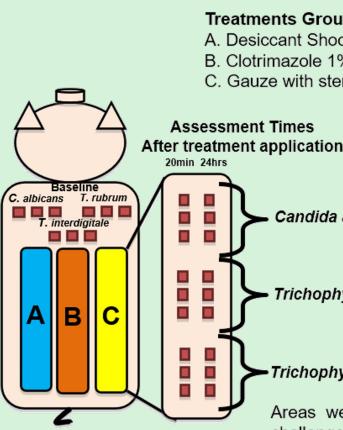
Swine were used as our experimental animal due to the morphological, physiological, and biochemical similarities between porcine skin and human skin.⁸

2. Wounding Technique:

A specialized electrokeratome was used to create Sixty-three (63) deep partial thickness wounds measuring (10 mm x 7 mm x 0.5 mm deep) on the paravertebral and thoracic area. 3. Inoculation:

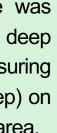
After creation of wounds, 100µl of Trichophyton rubrum ATCC28188 (TR28188), Trichophyton interdigitale ATCC9533 (TR9533) and Candida albicans ATCC64550 (CA64550) was used to inoculate each wound by scrubbing 10⁶ CFU/ml) inoculums into each wound with a teflon spatula (30 seconds). Nine (18) wounds were assigned to each treatment group (3 groups total) and 3 wounds were used as a baseline All wounds were then covered with a polyurethane film for 72 hours (to allow colonization).

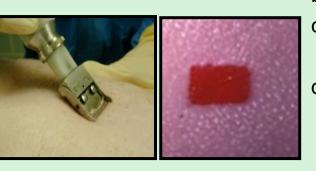
4. Experimental Design:



- · Serial dilutions (photo i) were made from all culture samples and the extent of Trichophyton interdigitale ATCC9533 microbiological contamination assessed using the Spiral Plater System (Spiral Biotech, Areas were uncovered after 3 days after Norwood, MA – photo j). This system deposits a 50µL aliquot of the scrub bacterial challenge. 4 infected wounds per organism were recovered as a baseline suspension over the surface of a rotating agar plate. BBL[™] CHROMagar[™] Candida was used to isolate CA64550 (photo k) and Dermatophyte Test Medium (photo I) was used to isolate the other 2 dermatophytes (TR28188 and TI9533). All plates were incubated References aerobically (24 hours – 5 days) at 30oC, after which the number of viable colonies were 1. Palackic A, Popp D, Tapking C, Houschyar KS, Branski LK. Fungal infection in Burn Patients. Surg Infect, 2021, 22(1):83-87. counted 2. Gnat S, Lagowski D, Nowakiewicz. Major challenges and perspectives in the diagnostics and treatment of dermatophyte
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Treatments Groups

A. Desiccant Shock Technology [DST*] B. Clotrimazole 1% Positive Control+ C. Gauze with sterile saline

- Candida albicans ATCC64550
- Trichophyton rubrum ATCC28188

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5. Treatment Regimen:

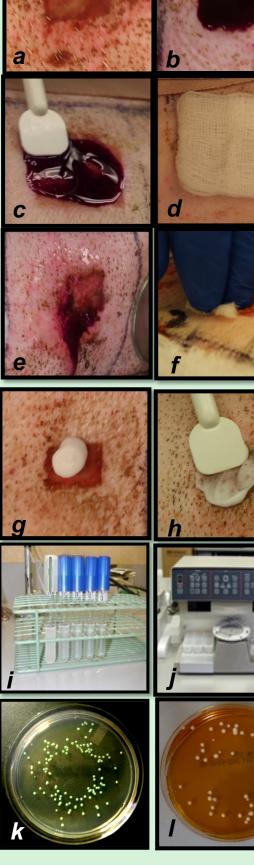
- a. After 72 hours, all wounds were treated
- b. Wounds treated with DST received 200ul
- DST treatment was spread with spatula and allowed to stay in place for 30 seconds
- Saline Irrigation wounds each had a premoisten gauze (200 µL of sterile saline) placed over the wound which was allowed to stay in place for 30 seconds.
- e. After 30 seconds, all wounds were rinsed with a 5mL syringe of sterile saline (image showed rinsing after NDF application).
- After rinse wounds were gently wipe with moistened sterile PBS gauze and then covered with Tegaderm.
- Clotrimazole 1% Positive Control wounds received 200mg of treated
- Positive control was spread with sterile spatula.

6. Wound Recovery:

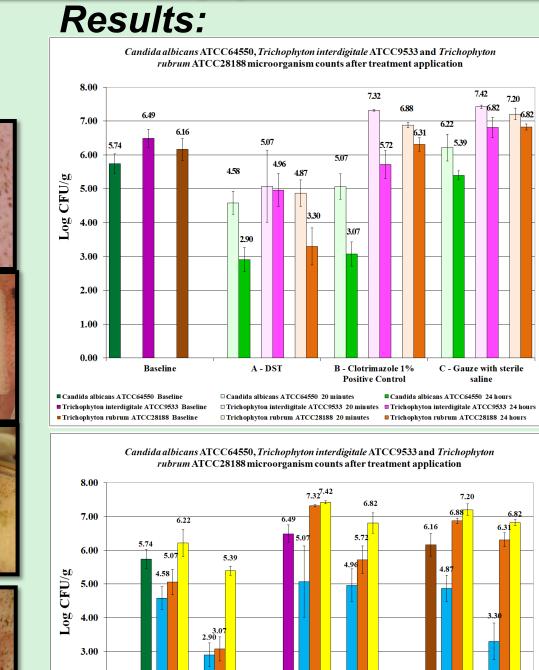
- On Day 0 (72 hours after inoculation), three wounds from each organism were biopsied (6mm punch) as a baseline. Then three treated wounds were biopsied (6mm punch biopsy) 20 minutes after treatment application for each treatment group. The remaining wounds were cultured at 24 hours after treatment application
- The biopsies (6mm) were weighed and immediately placed in 1 mL of All Purpose Neutralizing Solution. The sample was combined with an additional 4 mL of Neutralizing Solution and homogenized in a sterile homogenization tube.

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Candida albicans ATCC64550, Trichophyton interdigitale ATCC9533 and Trichophyton **4.00** Candida albicans Trichophyton interdigitale Trichophyton rubrum ATCC64550 ATCC9533 ATCC28188 Baseline
A – DST 24 hours
B - Clotrimazole 1% Positive Control 24 hours
C - Gauze with sterile saline 20 minutes

• When comparing baseline wounds results, those wound infected TI9533 exhibited the highest microorganism counts at 6.49 ± 0.27 Log CFU/g. Both CA64550 and TR28188 exhibited baseline counts of 5.74 \pm 0.29 and 6.16 ± 0.33 Log CFU/g, respectively.

- Those wounds treated with Gauze with Sterile Saline exhibited the highest fungal counts among the entire study for every microorganism.
- Wounds infected with CA64550 and TR28188 then treated with DST were able to reduce their fungal counts when comparing 20 minutes and 24 hours after treatment application.

• DST was the only treatment group that resulted in lower counts than baseline results (both 20 minutes and 24 hours).

 Fungal counts were exhibited after 24 hours with DST treated wounds showing fungal reductions of 97.06% and 99.86% when compared against TI9533 and TR28188, respectively. When challenging wounds treated with CA64550, both DST and Clotrimazole 1% Positive Control exhibited large fungal differences after 24 hours at 99.85% and 99.78%, respectively.

 For CA64550, after 24 hours, those wounds treated with DST had slightly lower fungal counts than wounds treated with Clotrimazole 1% Lamisil Positive Control at 2.90 \pm 0.36 and 3.07 ± 0.36 Log CFU/g, respectively.

 DST treated wounds, in both 20 minutes and 24 hours results, exhibited substantially lower fungal counts against CA64550 and TI9533 when compared against results of wounds infected with TR28188.

 DST showed compared against Gauze with Saline Control after 24 hours a fungal reduction of 98.60%. For those wounds infected with TR28188 after 20 minutes, DST treated wounds had substantially lower fungal counts, with fungal reductions of 99.02% and 99.54% when compared against Positive Control and Gauze with Saline

Conclusions

C - Gauze with sterile saline 24 hours

• Overall, those wounds treated with DST showed substantially lower fungal counts against the three microorganisms and in both 20 minutes and 24 hours. Clotrimazole 1% Positive Control had lower fungal counts when treating wounds infected with Candida albicans ATCC64550, however it did not appear as effective in wounds infected with either Trichophyton interdigitale ATCC9533 or Trichophyton rubrum ATCC28188. Additional samples would be required to substantiate these claims.