Frictional Fabric-Based Tissue Biopsy Sampling for Wound Organism Analysis

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SoftBiopsy®

Inspect the

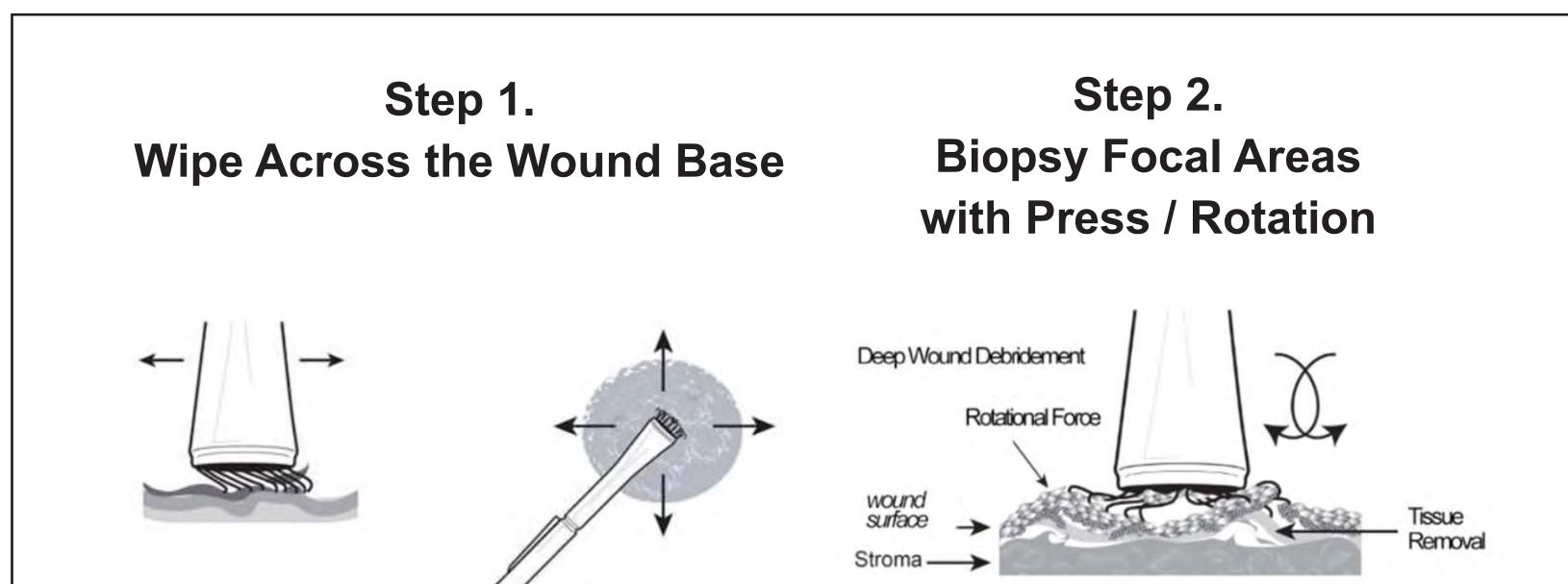
for tissue

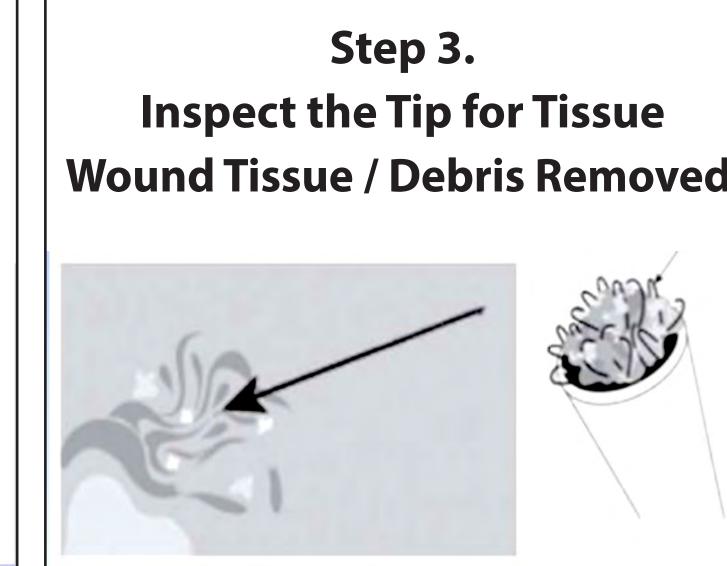
Introduction:

Evaluation of chronic non-healing wounds for the presence of planktonic and diagnostically sample wounds. Because traditional full thickness biopsy has been considered traumatic and aggressive, traditional Z-Swab samples of tissue fluids hasaided in removing sample fluid and cytology for anatomic, microbiologic, or molecular testing. This abstract introduces a new technology designed to be minimally invasive yet remove histologic biopsy samples suitable for analysis, which has been proven over 1.5 million clinical cases of cervical biopsies to gently remove trans-epithelial biopsies from intact lesion-bearing mucosal tissues.

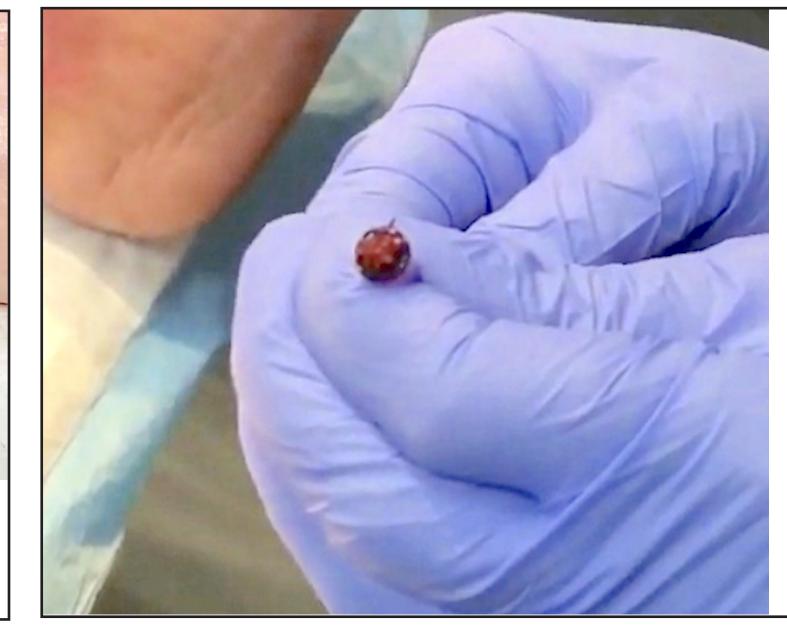
Methods:

After informed consent, the FDA compliant frictional fabric-based biopsy device SoftBiopsy® was used in post-debridement cases to sample the wound base and evaluated microscopically for evidence of intact tissue suitable for molecular, culture, or anatomic pathology testing. The method was to gently sweep the hooked fabric bristles across the entire wound surface, and secondarily focus on areas of apparent infection or possible biofilm, pressing and twisting the fabric as to frictionally abrade and capture curetting type fragments of tissue and trap them into the hooked fabric tip is then detached from the handle and sent to the lab for analysis.

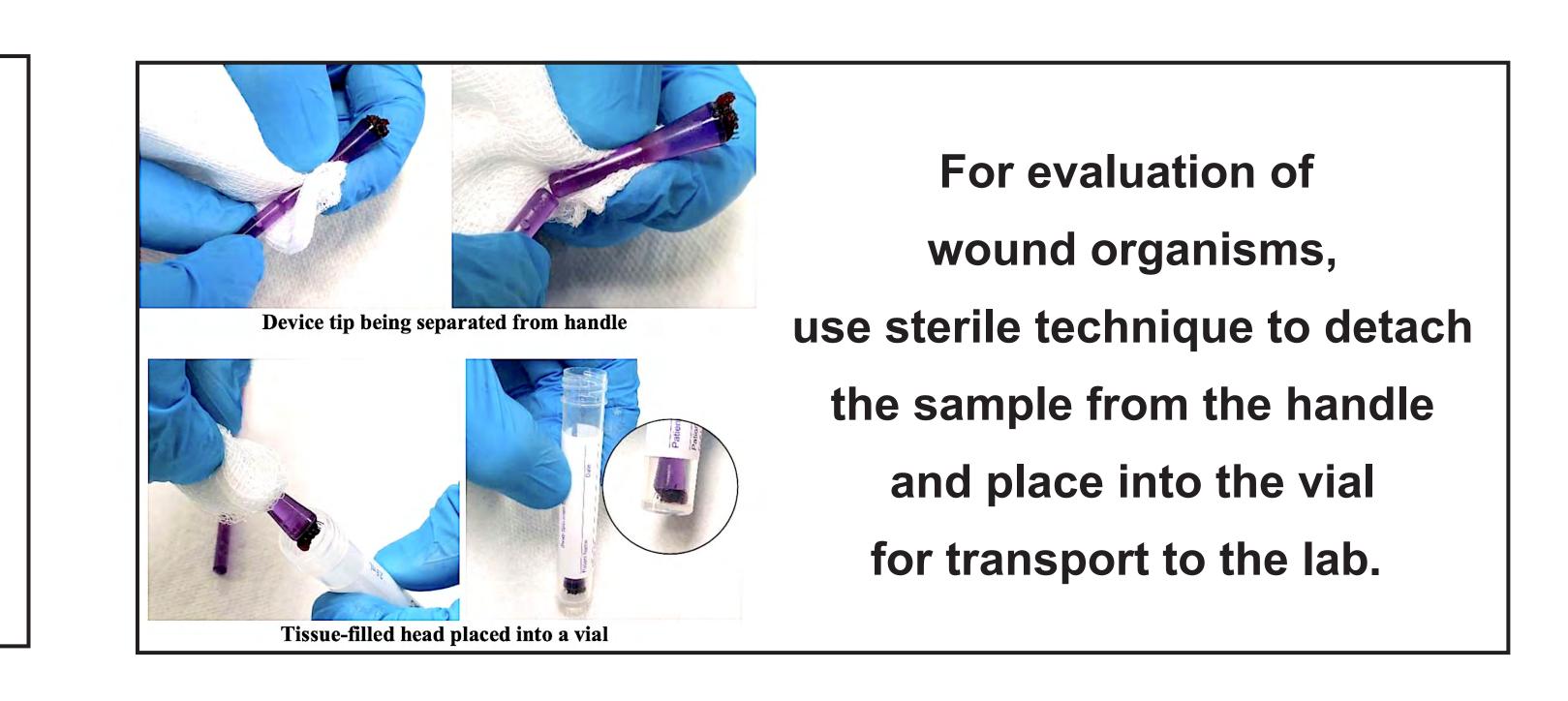






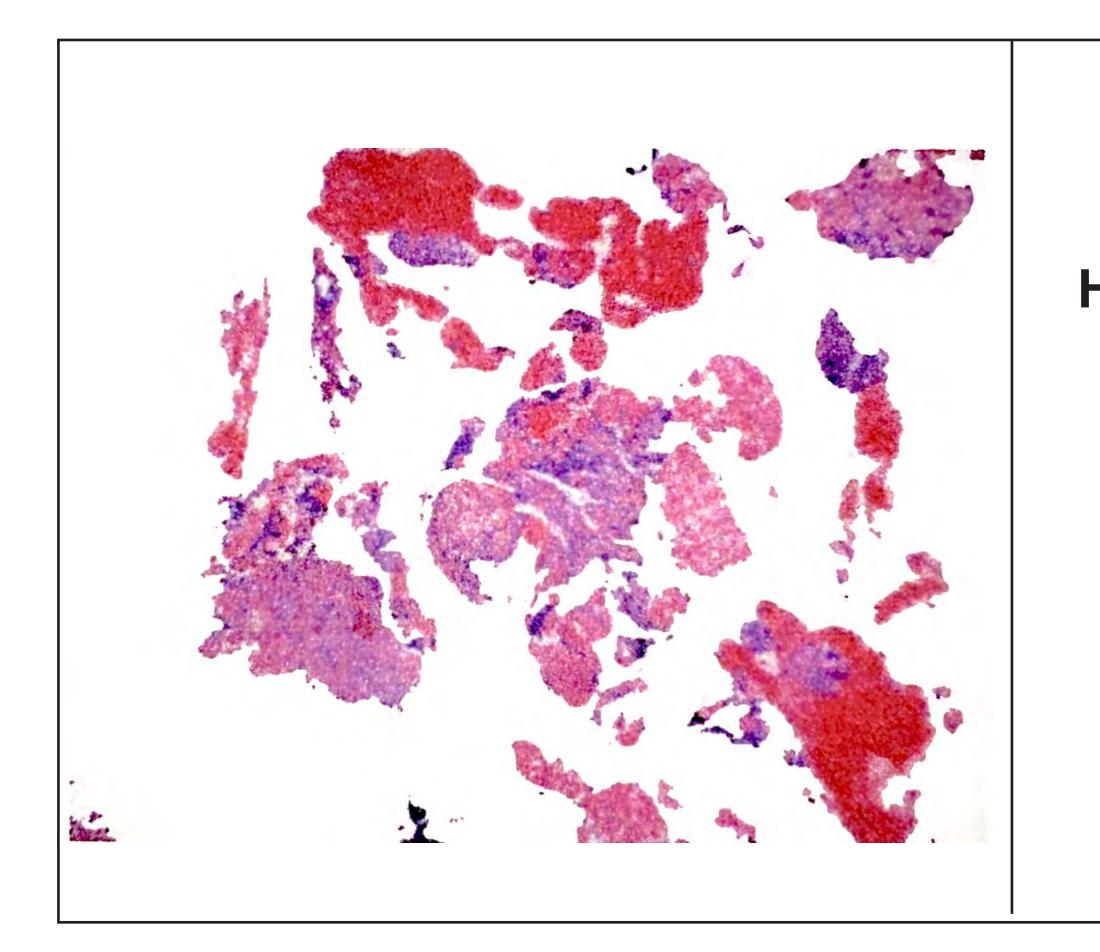


After Gently Removing the SoftBiosy® from the wound site, separate head (with Kylon® Pad and Tissue Sample) from body of device by snapping at the recessed joint Kylon® fabric after sampling Gløves should be worn



Results:

Photomicrographic evidence of tissue samples used for testing is presented. The acquisition of tissue curettage samples was perceived as minimally invasive by both the patient and clinician evaluating the patient during the tissue sampling.



Histopathologic evidence of tangential (curettage) biopsy results from SoftBiopsy® sampling of the debrided wound base.

RESULTS SUMMARY			
Organism Classification	Genus Species	Result	Microbial Load
AEROBIC BACTERIA: GRAM NEGATIVE	Klebsiella pneumoniae	DETECTED(+)	HIGH
AEROBIC BACTERIA: GRAM POSITIVE	Staphylococcus aureus	DETECTED(+)	HIGH
AEROBIC BACTERIA: GRAM POSITIVE	Enterococcus faecalis	DETECTED(+)	MEDIUM
AEROBIC BACTERIA: GRAM NEGATIVE	Pseudomonas aeruginosa	DETECTED(+)	LOW
			2
AEROBIC BACTERIA: GRAM POSITIVE	Corynebacterium jeikeium	DETECTED(+)	LOW
	Corynebacterium jeikeium Candida parapsilosis	DETECTED(+)	LOW
FUNGI Resistance Gene Classification	Candida parapsilosis Resistance Gene	DETECTED(+) Result	
FUNGI Resistance Gene Classification Extended Spectrum ß-lactamase	Candida parapsilosis Resistance Gene SHV	DETECTED(+) Result DETECTED(+)	
FUNGI Resistance Gene Classification Extended Spectrum ß-lactamase Extended Spectrum ß-lactamase	Candida parapsilosis Resistance Gene SHV TEM	DETECTED(+) Result DETECTED(+) DETECTED(+)	
FUNGI Resistance Gene Classification Extended Spectrum ß-lactamase Extended Spectrum ß-lactamase MRSA / MRSE potential	Candida parapsilosis Resistance Gene SHV TEM MecA	DETECTED(+) Result DETECTED(+) DETECTED(+) DETECTED(+)	
Resistance Gene Classification Extended Spectrum ß-lactamase Extended Spectrum ß-lactamase MRSA / MRSE potential Macrolide-Lincosamide-Streptogram B	Candida parapsilosis Resistance Gene SHV TEM MecA ErmA	DETECTED(+) Result DETECTED(+) DETECTED(+) DETECTED(+) DETECTED(+) DETECTED(+)	
AEROBIC BACTERIA: GRAM POSITIVE FUNGI Resistance Gene Classification Extended Spectrum ß-lactamase Extended Spectrum ß-lactamase MRSA / MRSE potential Macrolide-Lincosamide-Streptogram B Macrolide-Lincosamide-Streptogram B Tetracycline	Candida parapsilosis Resistance Gene SHV TEM MecA	DETECTED(+) Result DETECTED(+) DETECTED(+) DETECTED(+)	

Discussion:

Excavating areas suspect of biofilm in wounds using biopsy rather than traditional swabbing has been shown to present a robust sample for testing of organisms and antibiotic sensitivity. Frictional abrasion as a means for biopsy collection has been shown in mucosal tissue to be immune stimulatory. More research needs to be done in chronic wounds to evaluate this effect on ameliorating biofilm and enhancing wound repair.

References

- . Winter M. et al. Fabric-based exocervical and endocervical biopsy in comparison with punch biopsy and sharp curettage. J.Low Genit Tract Dis, 2012, 16(2): 80-7.
- 2. Diedrich J, Rathore S, Bentz J. Comparison of Tissue Yield Using Fricdtional Fabric Brush versus Sharp Curettage of Endocervical Curettage. J Low Genit Tract Dis, 2017, 21(4), 304-6.
- B. Sitelman A, Diedrich J, Lonky NM. Observation of a Robust Immune Inflammatory Response Following Frictional Fabric Cervical Biopsy and Endocervical Curettage During Colposcopy. J repro Med, 2019, 64(4), 261-4.
- 4. Bowler PG, Duerden BI, Armstrong DG. Wound microbiology and associated approaches to wound management. Clin Microbio Rev, April 2001, p. 244-269.
- 5. Copeland-Halperin LR, Kaminsky AJ, Bluefield N, Miraliakbari R. Sample procurement for cultures of infected wounds: a systematic review. J Wound Care, North American Supplement Vol25 (4), April 2016, p. S4-S10.
- 6. Huang Y, Cao Y, Zou M, et al. A Comparison of Tissue versus Swab Culturing of Infected Diabetic Foot Wounds. Int J Endocrinology, Volume 2016, Article ID 8198714, p1-6.
- 7. Pallua N, Fuchs PC, Hafemann B, et al. A new technique for quantitative bacterial assessment on burn wounds by modified dermabrasion. Journal of Hospital Infection (1999) 42: 329–337.
- 8. Melendez JH, Frankel YM, A. T. An AT, et al. Real-time PCR assays compared to culture-based approaches for identification of aerobic bacteria in chronic wounds. Clinical Microbiology and Infection, Volume 16 Number 12, December 2010, p 1762-69.
- 9. Rondas A, Schols JM, Ruud J.G. et al. Swabs versus biopsy for the diagnosis of chronic infected wounds. ADVANCES IN SKIN & WOUND CARE, MAY 2013, p 211-219.
- 10. Attinger C, Wolcott R. Clinically addressing biofilm in chronic wounds. ADVANCES IN WOUND CARE, VOLUME 1, NUMBER 3, 2012, p 127-132.



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