# Research Letter

# From the Cochrane Library: Hydrosurgical Debridement Versus Conventional Surgical Debridement for Acute Partial-Thickness Burns

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#### **KEYWORDS**

Cochrane; systematic review; randomized controlled trial; hydrosurgery; hydrosurgical debridement; debridement; burns; wound healing

Partial-thickness burns often require surgical excision with dressings or reconstruction. Standard of care includes early debridement (tangential excision of nonviable tissue) followed by split-thickness skin grafting. The goal of debridement is to reach a plane of viable tissue, while sparing healthy, uninjured tissue, expediting healing and minimizing scarring. Conventional debridement (scalpel or knife) is potentially limited by inaccurate differentiation between viable and nonviable tissues, with resultant delayed healing and greater scarring. Hydrosurgery is an alternative tool for surgical debridement that uses pressurized saline and a vacuum system to create a Venturi effect, ideally improving debridement accuracy and tissue-sparing. The Cochrane systematic review "Hydrosurgical debridement versus conventional surgical debridement for acute partial-thickness burns" analyzed existing randomized controlled trials (RCTs) enrolling participants with acute partial-thickness burn injuries requiring debridement and grafting; this yielded one eligible study randomizing 61 pediatric patients to either conventional debridement (n=31) or hydrosurgery (n=30) [1].

In this RCT, no clear differences were observed in the mean time to complete healing (mean difference [MD] 0 days, 95% CI –6.25 to 6.25), postoperative infection risk (risk ratio 1.33, 95% CI 0.57-3.11), operative time (MD 0.2 minutes, 95% CI –12.2 to 12.6), or 6-month scar outcome (MD not computed).

Study conclusions were very low certainty on the GRADE (Grading of Recommendations Assessment, Development and Evaluation) assessment, showed a high risk of reporting bias, and were limited by the small sample size (not powered to detect differences in primary outcomes). Generalizability was limited, as the study focused on a pediatric population and smaller burn injuries (3%-4% of total body surface area). No information was reported on clinical resource use, health-related quality of life, or adverse events. The authors concluded that it remains unknown if hydrosurgery is superior to conventional surgery for treatment of middepth burns.

Following the publication of the Cochrane review, no further RCTs have been published that compare the efficacy of hydrosurgical debridement to conventional blade debridement for burns. However, one study is still "awaiting classification," and one multicenter RCT (n=137) is underway to examine long-term (12 months) scar quality for hydrosurgical versus conventional debridement of dermal burns [2].

In addition to its application for burns, there is evidence for hydrosurgery treating other dermatological pathologies. For example, in a study of axillary osmidrosis (n=93), hydrosurgery showed improved patient satisfaction and fewer postoperative complications compared to traditional surgery [3]. Case reports of severe phymatous rosacea, which currently lacks standard

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surgical guidelines, document successful treatment with hydrosurgery [4]. Additionally, hydrosurgery can safely and rapidly debride various ulcer types in outpatient settings [5]. As the COVID-19 pandemic continues to decrease the availability of inpatient rooms and services, the possibility of providing outpatient hydrosurgical debridement for wounds may be important for continuing patient care. Dermatologists manage numerous wounds in daily practice; therefore, providers should be informed of the current recommendations for wound debridement. Future research should include additional high-quality RCTs comparing the efficacy of hydrosurgery versus standard debridement for burns. Outcome measures could focus on patient-reported scarring and adverse events. This would increase the certainty and generalizability of the results, and provide evidence for procedural recommendations.

### **Conflicts of Interest**

MDS is a member of the Cochrane Collaboration. TES is a member of the Cochrane Collaboration and an Editorial Board Member-at-Large for *JMIR Dermatology*, and receives fellowship funding from the Pfizer Global Medical Grant (58858477) Dermatology Fellowship 2020 (principal investigator RPD). RPD is a Joint Coordinating Editor for *Cochrane Skin*, a dermatology section editor for *UpToDate*, a Social Media Editor for the *Journal of the American Academy of Dermatology*, a Podcast Editor for the *Journal of Investigative Dermatology (JID)*, Editor-in-Chief of *JMIR Dermatology*, a coordinating editor representative on Cochrane Council, and Cochrane Council cochair and director of the University of Colorado Anschutz Medical Campus United States Cochrane Affiliate. RPD receives editorial stipends (*JMIR Dermatology, JID*), royalties (*UpToDate*), and expense reimbursement from Cochrane Skin. JCRW, NIHR Doctoral Research Fellow (NIHR301793), is funded by the National Institute for Health Research (NIHR) for this research project. The views expressed in this publication are those of the authors and not necessarily those of the NIHR, National Health Services, or the UK Department of Health and Social Care.

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#### Abbreviations

GRADE: Grading of Recommendations Assessment, Development and EvaluationJID: Journal of Investigative DermatologyMD: mean differenceNIHR: National Institute of Health ResearchRCT: randomized controlled trial

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