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Introduction and Background

- Preoperative infection management is crucial for reducing complications in chronic complex wound reconstruction, yet this often requires a multidisciplinary approach.
- Even extremely complex patients can be safely taken to reconstruction when infection and bacterial bioburden are managed objectively.
- Traditional infection assessment methods (e.g. wound swabs and biopsies) can take days to acquire results, thereby delaying intervention.
- Fluorescence imaging enables real-time, objective bacterial detection for faster and targeted infection control, potentially enhancing presurgical preparation, improving outcomes and shortening time to reconstruction [1-3].

This case series aims to show how fluorescence imaging can improve pre-operative infection control, wound bed preparation, and clinical decision marking ahead of complex, multidisciplinary surgical reconstructions via skin substitute application.

Material and Methods

- A fluorescence imaging device (MolecuLight i:X) was utilized pre-, intra- and post-operatively during complex wound cases involving surgical debridement, reconstruction, and skin substitute placement.
- Positive fluorescence indicated the presence of pathogenic bacterial loads >10⁴ CFU/g. Red fluorescence indicates Gram +/- bacteria and cyan fluorescence indicates *Pseudomonas aeruginosa*, while skin/slough fluoresce green.

Conclusions

- Pre-operative fluorescence imaging optimized wound bed preparation in the hospital inpatient setting, expediting surgical readiness and improving outcomes in complex skin substitute placement.
- Fluorescence imaging was involved in key decision points, including:



[1] Rahma S., Woods, J., Brown, S., et al. The Use of Point-of-Care Bacterial Autofluorescence Imaging in the Management of Diabetic Foot Ulcers: A Pilot Randomized Controlled Trial. Diabetes Care, 2022. Jul 7;45(7):1601-1609. [2] Kelso, M.K., and Jaros, M. Improving Wound Healing and Infection Control in Long-term Care with Bacterial Fluorescence Imaging. Adv Skin Wound Care, 2024. Sep 1;37(9):471-479. [3] Price N. Routine Fluorescence Imaging to Detect Wound Bacteria Reduces Antibiotic Use and Antimicrobial Dressing Expenditure While Improving Healing Rates: Retrospective Analysis of 229 Foot Ulcers. *Diagnostics*, 2020. Nov 10;10(11):927.

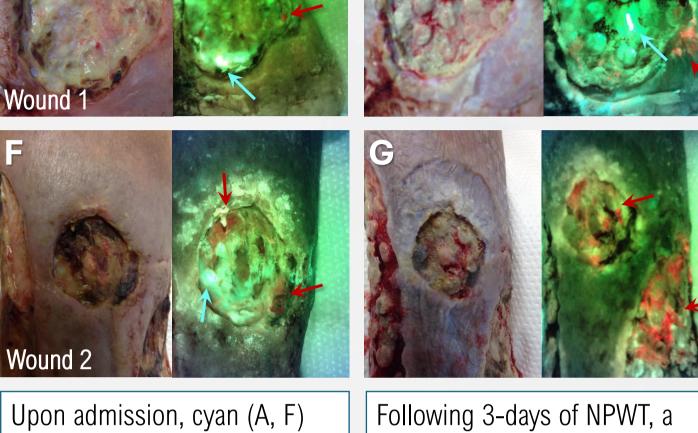
Pre-operative Wound Bed Preparation Using Fluorescence Imaging to Optimize Outcomes in Chronic Wound Reconstruction

Case 1 – Multiple Chronic Lower Extremity Ulcerations **Initial Visit** 3 Days 7 Days

Red/cyan fluorescence decreasing (C-D, H-I). I&D required, but popliteal artery aneurysm perforated. New NPWT with instillation was applied while waiting for vascular bypass.



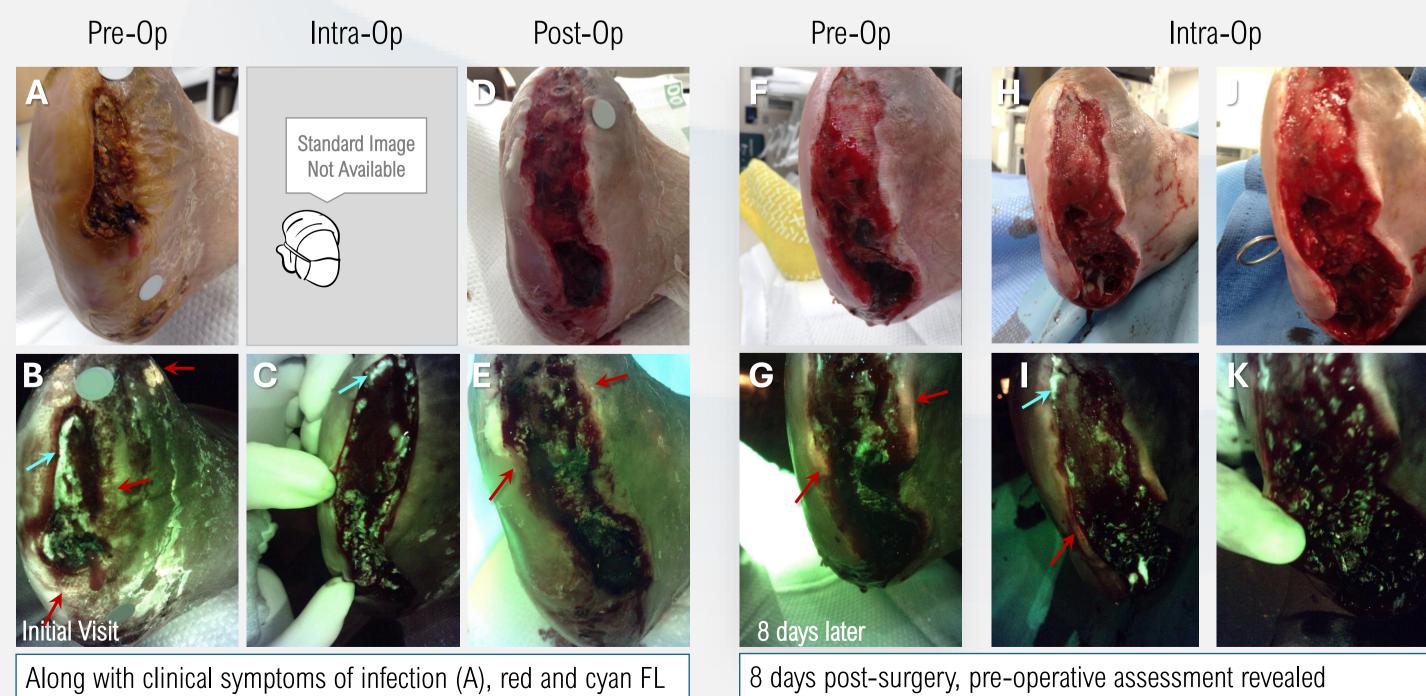
multidisciplinary



and red/yellow (A, F) fluorescence signals (arrows) were observed along with abundant slough and necrotic tissue debris

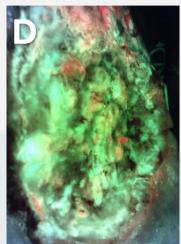
heavy amount of bioburden speculated, as fluorescence imaging uncovers persistent red and cyan signals. Wound VAC with instillation applied.

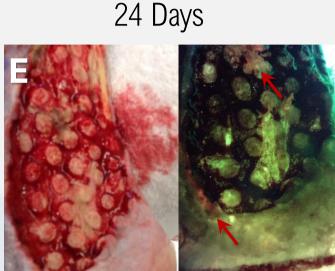
Case 2 – Infectious Complications Following a Transmetatarsal Amputation

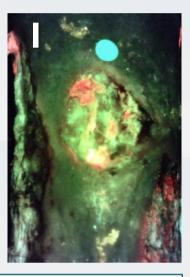


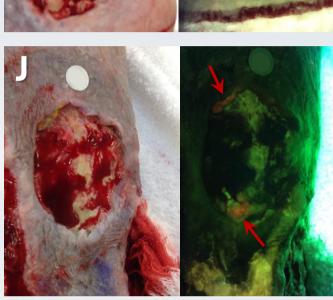
signals were pre-operatively observed on the wound bed and periwound (B). Intra-operative fluorescence-guided surgical debridement reduced FL signals (C). NPWT was applied postoperatively (D), yet red FL on the periwound persisted (E).

9 Days

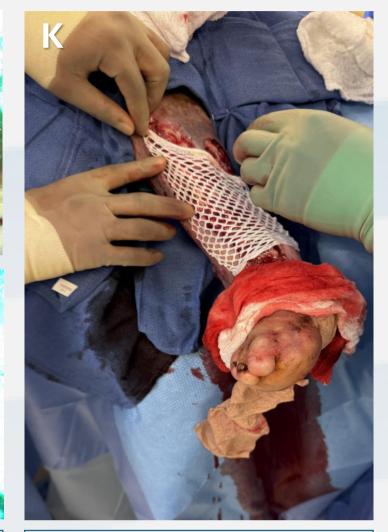








Post-bypass and revascularization, fluorescenceguided debridement significantly reduced bacterial burden as shown by decreased fluorescence signals (E,J).



Intra-Op

With bacterial burden minimized, SomaGen meshed allograft and Salera mini membranes were applied to all wounds.

- and signs of infection.



Case Summary: In a complex patient involving multiple lower extremity ulcerations, pre-operative fluorescence imaging identified persistent red and cyan signals indicative of bioburden/biofilm, guiding timing of wound reconstruction. Intra-operative fluorescence imaging guided targeted debridement, leading to improved wound bed preparation for CAMP application.



excessive bleeding and oozing (F) and persistent red and cyan FL signals (G). Intra-operative FL-guided debridement was conducted (H), but FL was not completely eliminated (I). Guided debridement continued until almost all FL was removed (J-K).

Post-Op



With bacterial burden minimized SomaGen meshed allograft and Salera mini membranes were applied.



Case Summary: Despite standard of care treatments and surgical debridements, this post-amputation wound continued to deteriorate. Pre-operative fluorescence imaging revealed persistent bioburden/biofilm that was likely overlooked with conventional methods. These fluorescence signals were effectively reduced with intra-operative fluorescence-targeted debridement, leading to optimal wound bed preparation for CAMP application.

• 63 y.o. male with a history of peripheral arterial disease, venous insufficiency with ulceration, hypertension, hyperlipidemia, and uncontrolled type 2 diabetes, and multiple recurrent ulcers requiring surgical debridement with reconstruction.

Presented with 5 active lower extremity ulcerations (>2 years) with exposed tendon on right lower extremity with pain, heavy slough/exudate

• FL imaging showed red and cyan FL signals, confirmed by wound culture for *P. aeruginosa* and heavy growth of mixed morphologies.

Requires vascular surgery to improve blood flow, NPWT with instillation, and reconstruction with skin substitutes.

• 64 y.o. female with a history of hypertension, hyperlipidemia, hypothyroidism, and type-2 diabetes underwent a transmetatarsal amputation and revascularization of the left foot secondary to a diabetic foot ulcer with abscess, gangrene and osteomyelitis.

Patient underwent weekly surgical debridement, HBOT, and NPWT, but developed pain, abscess, and cellulitis confirmed by wound culture for *P*. aeruginosa and Corynebacterium sp.

• IV antibiotics ordered but not administered due to resourcing and patient compliance issues. Left foot worsened with pain, erythema, and abscess. • Requires NPWT with instillation, and reconstruction with skin substitutes.