

## Introduction

Chronic and acute wounds require etiology-specific assessment to guide timely interventions.<sup>1</sup> While traditional macro-vascular assessments, such as the Ankle-Brachial Index (ABI), are standard practice, they often fail to capture localized micro-perfusion deficits or early inflammatory changes. Multimodal point-of-care imaging, including near-infrared spectroscopy (NIRS) for tissue oxygenation (StO<sub>2</sub>),<sup>2</sup> thermography for skin surface temperature, and digital photography, offers non-invasive, real-time insights into perfusion, ischemia, and surrogate markers of infection.

## Methods

A case series was conducted to evaluate the clinical utility of multimodal point-of-care imaging in patients presenting with various lower extremity wounds. Assessments were performed using a handheld device (MIMOSA Pro, MIMOSA Diagnostics Inc.) capable of capturing NIRS, thermography, and digital photography.

**NIRS**

Assessment of overall perfusion and localized StO<sub>2</sub> levels within the wound bed and periwound tissue.

**THERMOGRAPHY**

Measurement of skin surface temperature gradients to identify infection or inflammation.

**BILATERAL COMPARISON**

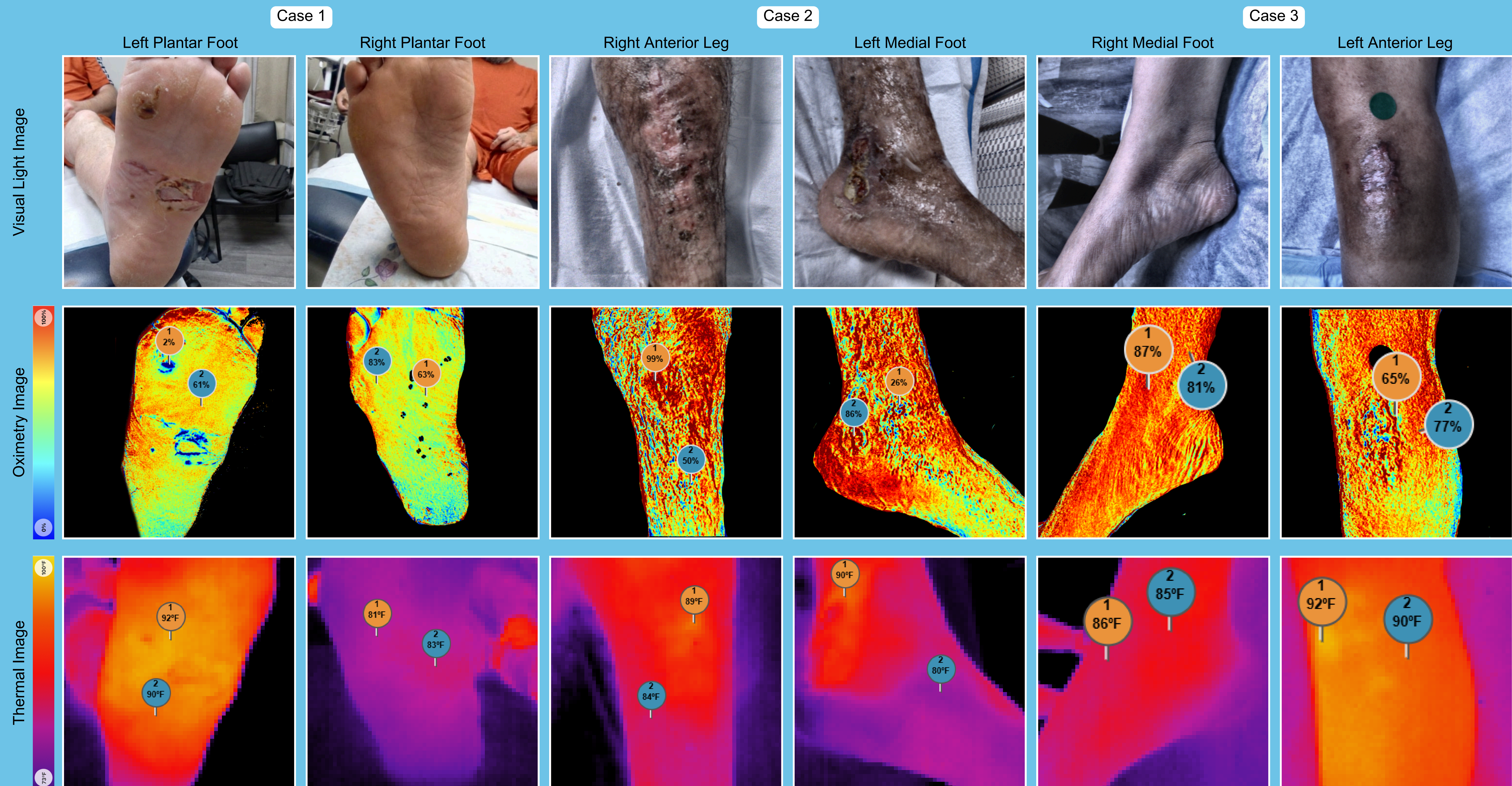
Comparative analysis of StO<sub>2</sub> and temperature between the affected limb and the contralateral limb to establish baseline values.

**WOUND BED**

Specific evaluation of localized StO<sub>2</sub> deficits (e.g., <39%) to determine the necessity of clinical interventions such as debridement.

Clinical Assessment Methodologies Using Multimodal Point-of-Care Imaging. StO<sub>2</sub>: Tissue Oxygen Saturation (or Tissue Oxygenation)

## Results



**Case 1 - diabetic foot ulcer (DFU):** A 46-year-old male with type 2 diabetes presented with a left plantar DFU and normal bilateral ABIs (1.2). NIRS confirmed sufficient periwound perfusion with bilateral StO<sub>2</sub> > 50%, but identified a localized wound bed deficit (StO<sub>2</sub> < 39%) indicating a need for debridement. Thermography showed minor elevation in the wound bed (92°F) versus surrounding tissue (90–91°F), indicating no active infection.

**Case 2 - pyoderma gangrenosum (PG):** A male patient presented with PG on the right anterior leg and medial foot with active infection and normal ABIs (1.3). NIRS showed hyperperfusion (StO<sub>2</sub> > 70%) overall, but identified localized ischemic zones (StO<sub>2</sub> down to 0%). Thermal imaging confirmed infection via significant gradients exceeding 4°F.

**Case 3 - burn:** An 84-year-old female presented with a traumatic burn on the left anterior leg and normal bilateral ABIs. NIRS imaging showed overall StO<sub>2</sub> > 50%, while small areas of decreased perfusion (StO<sub>2</sub> < 39%) were detected in the wound bed, they did not necessitate debridement. Thermography showed no significant gradients (>2°F), suggesting an absence of infection.

ABI: Ankle-Brachial Index; DFU: Diabetic Foot Ulcer; NIRS: Near-Infrared Spectroscopy; PG: Pyoderma Gangrenosum; PVD: Peripheral Vascular Disease; StO<sub>2</sub>: Tissue Oxygen Saturation (or Tissue Oxygenation)

## Discussion and Key Takeaways

Multimodal point-of-care imaging facilitates etiology-specific assessment for chronic and acute wounds by integrating perfusion data and thermography as a surrogate for infection, thereby enhancing clinical decision-making. By combining NIRS and thermography, clinicians can objectively evaluate wound bed preparation requirements, such as identifying the specific need for debridement in areas of StO<sub>2</sub> < 39% or, conversely, avoiding unnecessary intervention. This approach confirms perfusion adequacy to support healing across diverse conditions—including diabetic foot ulcers, pyoderma gangrenosum, and burns—even when macro-perfusion markers like ABI appear normal. Furthermore, the integration of temperature gradients allows for the real-time detection of infection by identifying localized heat elevations (e.g., >4°F) relative to surrounding tissue or the contralateral limb. Ultimately, this multimodal framework enables a more precise, data-driven strategy for managing complex wound etiologies.

1. Kelso, M. R. et al. Use of near infrared spectroscopy in post-acute care: analysis of real-world clinical decision-making. J Wound Care 34, S6–S14 (2025).  
 2. Oropallo, A. et al. Advancing chronic wound care with near-infrared spectroscopy imaging: clinical applications, measurement parameters, and insights into healing dynamics. Wounds 37, 384–392 (2025).