Clinical Assessment of Protopic® Ointment in Deep Partial-Thickness Burns

NCT05856994

Status	RECRUITING
Phase	Phase 1
Sponsor	Vanderbilt University Medical Center
Enrollment	18 participants

Key Eligibility Criteria

Inclusion (7)

- Provision of signed and dated informed consent form by the subject or Legally Authorized Representative.
- Stated willingness to comply with all study procedures and availability for the duration of the study.
- Male or female aged 50 to 75 at time of screening visit.
- For females of reproductive potential, confirmed negative urine pregnancy test at enrollment.
- Presence of deep partial-thickness burns to one or both dorsal hands (burns may be present on other areas of the body, so long as all regions combined are d 5% Total Body Surface Area \[TBSA\], using the Browder and Lund Chart).

... and 2 more (see full listing online)

Exclusion (13)

- Pregnant, breastfeeding, or unwilling to practice birth control during participation in the study, if applicable.
- Presence of a condition or abnormality that in the opinion of the Investigator would compromise the safety of the patient or the quality of the data.
- Allergy or hypersensitivity to tacrolimus or other components of the ointment (per subject report) or personal preference.
- Allergy or hypersensitivity to bacitracin and/or any of its drug formulation components. Patients with known hypersensitivity to neomycin may also be sensitive to bacitracin.
- Subject is incarcerated.

... and 8 more (see full listing online)

Locations (1 total)

Vanderbilt University Medical Center, Nashville, Tennessee, United States

DISCLAIMER: This document is for informational purposes only and does not constitute medical advice. Always consult your healthcare provider before enrolling in any clinical trial. Information may not be up to date — verify details at ClinicalTrials.gov. Generated by ClinicalTrialsFinder.org.