

Early Intervention with the geko™ device for Venous Leg Ulcers Predicted to Not Heal Within 24 Weeks



Authors: Rabley-Koch C. A., Duong R., Cadavez R., Attard J., Entredicho A., Oshalla H., Orr A., Ramage D., Burrows C.

Aim

Quality Improvement Initiative aimed at enhancing policies and procedures so that patients can receive the most timely and effective practice to ensure or achieve better outcomes.

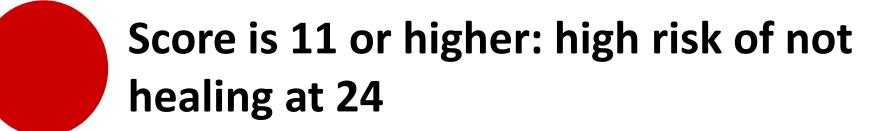
Procedure/Method

Earlier use of the geko™ device was employed on patients whose venous leg ulcers (VLU) that had an elevated risk of failure to close within 24 weeks. Patients were admitted to community clinic settings at 2 Ontario nursing agencies. Eleven patients were assessed twice over two weeks using a Validated Leg Ulcer Risk Assessment tool (VLURA). Moderate to high risk scores had geko™ devices added to their standard of care for a maximum of 12 weeks. Low scores were reassessed in two weeks; those increasing to moderate were started on the geko™ device.



Validated Leg Ulcer Risk Assessment Tool

(adapted)



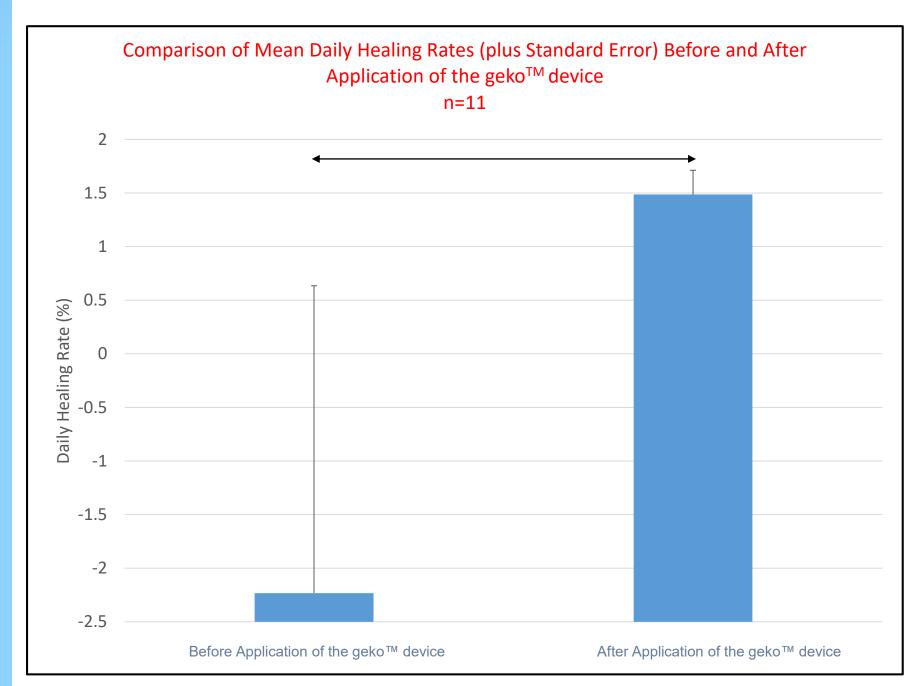




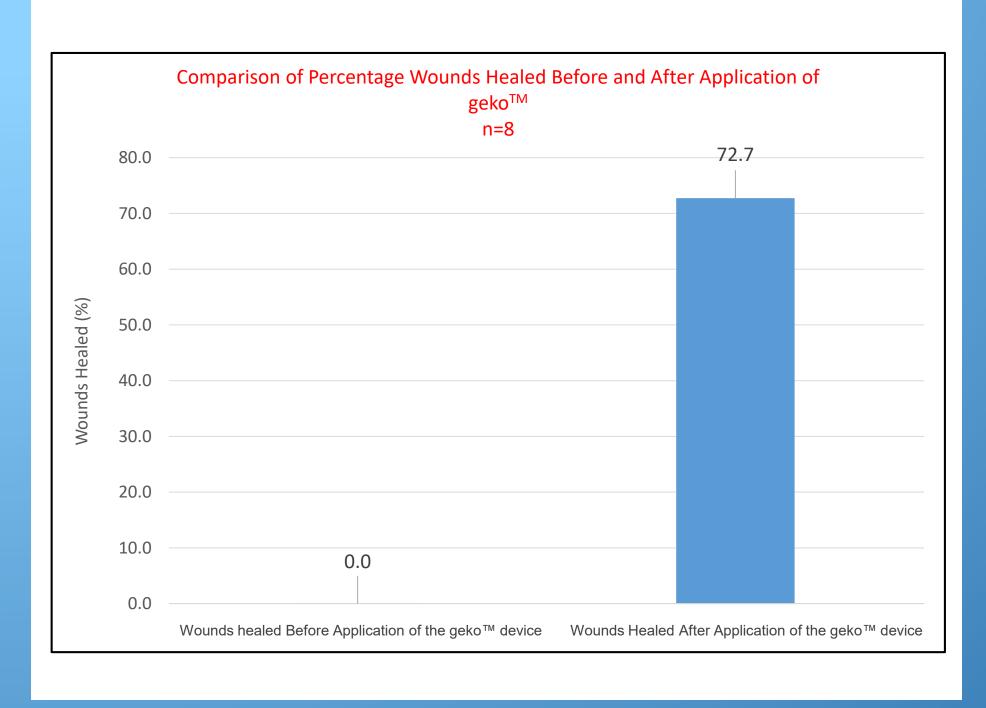
Findings/Results

- Frequent delays in the geko™ device initiation were related to a routine LHIN policy need to access vascular studies
- An average of 48.9 days elapsed between admission to device application
- Ten out of eleven patients experienced increased risk scores within the two weeks between initial and follow-up visit
- Preliminary results indicate a total of 12 wounds in 11 patients (80%) healed
- 2 wounds (13%) remained open with an average decrease in size of 88%
- One wound (6.7%) reopened
- Without use of the geko[™] device the average time for VLU closure in MH LHIN is 15 weeks
- Healing time with the geko[™] device is an average of 12 weeks

Healing Rates Before and After geko™ Application



Percentage Wounds Healed Before and After geko™ application



Implications/Applications

- Delays in access to timely care negatively impact wound healing
- Implementing a VLURA tool on admission identifies wounds with the greatest risk of failure to close
- Early intervention using the geko™
 device improves healing outcomes and
 decreases nursing visits
- Delay in the geko™ device initiation was related to clinician access to vascular studies/ABPI
- More work will need to be conducted to explore this further, particularly with the geko™ device application immediately upon referral

References

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