

Real-world Experience With a Decellularized Dehydrated Human Amniotic Membrane Allograft: A Prospective, Observational, Multicenter Study of a Broad Patient Population in All Wound Types. *Wounds* 2015;27(6):158-169

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The aim of this observational study was to gain experience in the use and performance of BIOVANCE[®] versus standard of care (SOC) in a real-world wound population. **A broad range of partial and full thickness wounds were studied across a mix of patient types.**

- Eligibility for inclusion included any patient that would benefit from treatment
- Unlike other chronic wound prospective, randomized, controlled trials, there were no limits on patients' age, baseline wound size or co-existing conditions
- Key comorbidities included: arterial insufficiency, autoimmune disease, diabetes, and edema/lymphedema

BIOVANCE SUPPORTS WOUND CLOSURE ACROSS A VARIETY OF WOUND AND PATIENT TYPES

BIOVANCE: at 8 weeks



Wounds < 3.12 cm



Wounds > 13.13 cm



SOC: at 12 weeks



(average baseline)



2X

BIOVANCE closed nearly twice as many wounds as control

The authors concluded that a longer observation time may have resulted in closure of the larger wounds.

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CASE STUDIES OF PATIENTS IN THIS STUDY

CASE STUDY 1: VENOUS STASIS ULCER

Patient	Comorbidities	Baseline Wound Size	Closure
<ul style="list-style-type: none"> 61 year old female with 3 prior treatment failures 	<ul style="list-style-type: none"> Peripheral vascular disease Venous insufficiency Immuno deficiency 	1.8cm x 1.2cm x 0.2cm	7 weeks with 1 application



BASELINE



WEEK 7 CLOSED

Treatment Regimen:

- Burow's Solution was applied to the periwound area twice a day

CASE STUDY 2: ACUTE WOUND

Patient	Comorbidities	Baseline Wound Size	Closure
<ul style="list-style-type: none"> 67 year old male with right BKA stump dehiscence 	<ul style="list-style-type: none"> Severe peripheral artery disease Diabetes 	7cm x 2.5cm x 0.1cm	6 weeks with 1 application



BASELINE



WEEK 6 CLOSED

Treatment Regimen:

- 1 application of BIOVANCE, not fenestrated
- Secondary dressing with nonadherent petroleum gauze and moist gauze

CASE STUDY 3: DIABETIC FOOT ULCER

Patient	Comorbidities	Baseline Wound Size	Closure
<ul style="list-style-type: none"> 68 year old male with full thickness wound 	<ul style="list-style-type: none"> Type 2 DM, Chronic Renal Failure, Neuropathic, Lymphedema 	12.9cm x 4.8cm x 0.1cm	25 weeks with 5 applications



BASELINE



WEEK 25 CLOSED

Treatment Regimen:

- Application of BIOVANCE on Days 1, 4, 16, 38 and 136
- Secondary dressing during treatment with petroleum gauze, topical gentamicin, silver hydrofiber dressing, polyurethane foam with gauze and elastic wrap
- Patient placed on gentamicin at week 3 with positive cultures for *staphylococcus aureus*

CITATION: Data and photos provided by Terry Treadwell, MD, on file at Celularity.

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