

USE OF PREVENA RESTOR BELLA-FORM™ INCISION MANAGEMENT SYSTEM IN BILATERAL, NIPPLE-SPARING MASTECTOMY

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PATIENT

A 42-year-old female presented to the surgical clinic after a prior diagnosis of invasive ductal carcinoma of the right breast (**Figure 1**). The patient presented with no concomitant comorbidities. Her medical history included asthma for which she had no recent use of her rescue inhaler or hospitalization.

PROCEDURE

The patient underwent bilateral, nipple-areolar-sparing mastectomy. The mastectomy produced two incisions at the inframammary folds (**Figure 2A**). Bilateral pre-pectoral direct-to-implant reconstruction was performed with round silicone implants (Natrella Inspira® SSX 340cc BI, model SSX-340; Allergan, plc, Dublin, Ireland). The implants were covered with an acellular dermal matrix. Bilateral drains were inserted and positioned laterally. Closure of the surgical incisions was achieved via subcuticular suturing (**Figure 2B**).

INITIAL TREATMENT/APPLICATION OF PREVENA RESTOR BELLA-FORM™ INCISION MANAGEMENT SYSTEM

PREVENA RESTOR BELLA-FORM™ Incision Management System (KCI, an ACELITY Company; San Antonio, Texas) was enlisted to help reduce tensile force across the closed incisions, manage the incision and surrounding area, and to help hold the edges of the closed incisions together. The PREVENA RESTOR BELLA-FORM™ Dressing (21cm x 19cm) was placed along the inframammary fold and mastectomy flap incisions (**Figure 3A**). The BELLA-FORM™ Dressing was positioned so as the drains exited outside of the perimeter of the dressing (**Figure 3B**). The PREVENA RESTOR™ Therapy unit applied -125mmHg of subatmospheric pressure for 6 days (**Figure 3C**).

DISCHARGE AND FOLLOW-UP:

On postoperative day (POD) 6, the patient returned to the clinic for a follow-up visit (**Figure 4A**). The PREVENA RESTOR BELLA-FORM™ System was removed to evaluate the reconstructed breasts, tissue integrity, and the incisions (**Figure 4B**). Upon dressing removal, some skin wrinkling was noted, which was temporary and did not cause any skin breakdown. Following the evaluation, the PREVENA RESTOR BELLA-FORM™ System was reapplied to manage the incision, reduce tensile force across the closed incision and to help hold the edges of the closed incision together. On POD 7, the patient returned to the clinic and the dressing was removed due to loss of seal as the patient was diaphoretic during the evening. Although erythema was present, this was not caused by the dressing (**Figure 5**). The patient was transitioned to a cotton brassiere with no pressure as well as prescribed topical silver sulfadiazine (Silvadene® Cream, Pfizer Inc., New York, NY) and oral ciprofloxacin. The drains were removed after 10 days. At POD 65, the incisions were completely healed, and both nipples were viable (**Figure 6**).



Figure 1. Patient at presentation after diagnosis of invasive ductal carcinoma of the right breast.

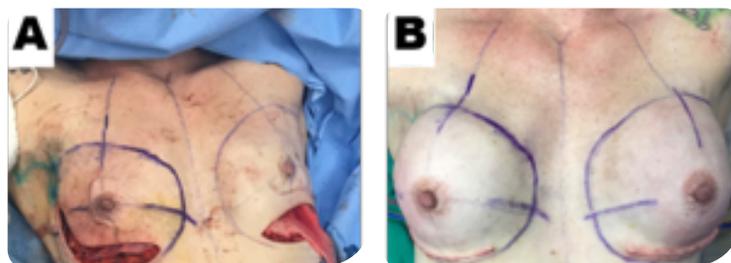


Figure 2. Bilateral, nipple-areolar-sparing mastectomy. **A.** Two surgical incisions at the inframammary folds resultant of bilateral mastectomy. **B.** Closure of bilateral incisions via subcuticular suturing.

USER EXPERIENCE

In this case, the PREVENA RESTOR BELLA-FORM™ System managed the incisions, helped hold the edges of the incision together, bolstered the mastectomy flap, and helped reduce skin edema. Nipple viability was maintained.

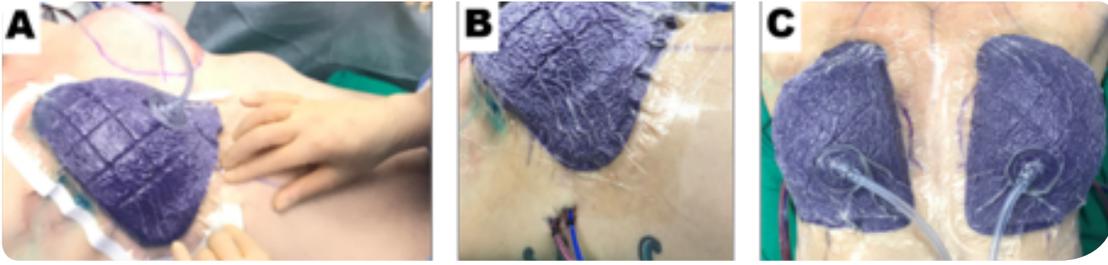


Figure 3. Application of PREVENA RESTOR BELLA-FORM™ Incision Management System. **A.** Placement of PREVENA RESTOR BELLA-FORM™ Dressing (21cm x 19cm) along the inframammary fold and mastectomy flap incisions. **B.** Dressing placement did not obstruct drains. **C.** The initiation of PREVENA™ RESTOR Therapy (-125mmHg) to the closed incisions.



Figure 4. POD 6 postoperative follow-up visit. BELLA-FORM™ System was reapplied. **A.** Patient before the removal of PREVENA RESTOR BELLA-FORM™ Dressing. **B.** Reconstructed breasts after removal of PREVENA RESTOR BELLA-FORM™ Dressing and before the reapplication of PREVENA™ Therapy.

Figure 5. On POD 7, erythema was noted bilaterally on the reconstructed breasts after removal of PREVENA RESTOR BELLA-FORM™ Dressing.



Figure 6. POD 65, incisions were completely healed, and both nipples were viable.

Photos and patient information courtesy of Dr. Regina M. Fearmonti, Alon Aesthetics/Fearmonti Plastic Surgery.

As with any case study, the results and outcomes should not be interpreted as a guarantee or warranty of similar results. Individual results may vary depending on the patient's circumstances and condition.

NOTE: Specific indications, contraindications, warnings, precautions and safety information exist for PREVENA™ Therapy. Please consult the applicable PREVENA™ System Clinician Guide instructions for use prior to application. Rx only.

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