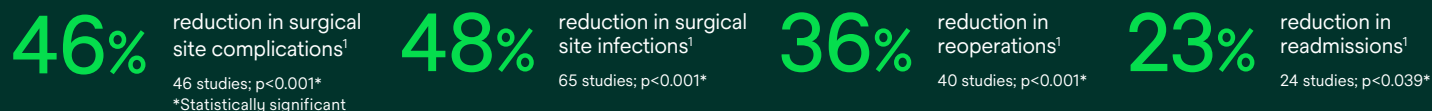


Your guide to help improve patient outcomes

A 2023 multi-specialty meta-analysis of 84 studies shows that Prevena Therapy has demonstrated significant improvement in key patient outcomes across multiple specialties.¹



Patient and procedure risk stratification backed by clinical evidence

While surgical patients may benefit from Prevena Therapy, patients at risk for complications such as surgical site infection may see added benefit. The following uses select study data²⁻¹⁸ to provide an illustrative guide to aid in risk stratification. This is not an all-inclusive list of risk factors. Clinicians are advised to use their clinical judgment to identify at-risk patients and procedures.

► Start here

Patient risk stratification

► Does the patient have **at least one** of the following risk factors for developing surgical site complications?

- | | | | |
|---|---|--|--|
| ☐ Active tobacco or nicotine use | ☐ Diabetes mellitus | ☐ Immunosuppression | ☐ Preoperative steroid or immunosuppressant use |
| ☐ Advanced age | ☐ Diffuse atherosclerotic disease | ☐ Malignancy | ☐ Prosthetic vascular graft |
| ☐ Autoimmune disease | ☐ Dyslipidemia | ☐ Malnutrition | ☐ Pulmonary disease |
| ☐ Chronic inflammatory disease | ☐ Elevated cardiac risk (hypertension, coronary heart disease, history of myocardial infarction) | ☐ Neoadjuvant chemotherapy | ☐ Significant pannus |
| ☐ Chronic kidney disease | ☐ Hypercholesterolemia | ☐ Non-aspirin anticoagulation | ☐ Staphylococcus aureus nasal colonization |
| ☐ Chronic obstructive pulmonary disease (COPD) | ☐ Hyperhomocysteinemia | ☐ Obesity (e.g., BMI>30 kg/m ²) | |
| ☐ Corticosteroid usage | | ☐ Peripheral vascular disease | |
| | | ☐ Preoperative bile stent/drain | |

Yes

No

Procedure risk stratification

► Is the procedure **high risk**?

- | | | | |
|---|--|---|--|
| ☐ Above knee amputation | ☐ High risk of edema | ☐ Nonlinear, intersecting, or branching incisions | ☐ Signs of hypoperfusion near the incision |
| ☐ After closure with flaps | ☐ High risk of lymphedema | ☐ Over new incisions that cross over previously healed incisions | ☐ Soilage risk |
| ☐ Ascites | ☐ High risk of seroma formation | ☐ Post-bariatric abdominoplasty | ☐ Sternotomy |
| ☐ Below knee amputation | ☐ High-tension closures/incisions | ☐ Pre or postoperative stoma | ☐ Synthetic graft implantations |
| ☐ Breast reconstruction | ☐ Large soft tissue defects | ☐ Preexisting wound infection | ☐ Thin soft tissue envelope |
| ☐ Contaminated or dirty wound | ☐ Large, undermined areas around the incision | ☐ Previous radiation therapy | ☐ Traumatized soft tissue |
| ☐ Emergency surgery | ☐ Multiple incisions | ☐ Repeated incisions | ☐ When delayed incision healing would postpone adjuvant therapy |
| ☐ Extended surgical time | ☐ Multiple stages of fracture repair | ☐ Revision surgery | |
| ☐ Extensive undermining | ☐ Need to reduce seroma space | ☐ Scarring concerns | |
| ☐ High energy trauma | | | |
| ☐ High risk of compromised perfusion | | | |

Yes

No

Consider Prevena Therapy

For additional safety information and instructions for use, contact your local Solventum representative.

Standard dressing

Versatility that meets your patients' needs

Solventum offers an expansive portfolio of negative pressure therapy dressings for closed incisions in several configurations and sizes to meet the needs of the surgeon while providing coverage for various surgical procedures and anatomical locations.

Protect the incision

Solventum™ Prevena™ Therapy Dressings

Manage the environment of closed incision via continuous NPWT at -125mm/Hg



Manage the incision and surrounding soft tissue

Solventum™ Prevena Restor™ Dressings

Manage the incision and surrounding soft tissue via continuous NPWT at -125mm/Hg with expanded coverage for the incision and surrounding soft tissue



Solventum™ Prevena™ 125 Therapy Unit

7-Day Unit



Solventum™ Prevena Plus™ 125 Therapy Units

7-Day Unit



14-Day Unit



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Solventum™ Prevena™ 125 Therapy Unit and Solventum™ Prevena Plus™ 125 Therapy Unit manage the environment of closed surgical incisions and remove fluid away from the surgical incision via the application of -125mmHg continuous negative pressure. When used with legally marketed compatible dressings, Prevena 125 and Prevena Plus 125 Therapy Units are intended to aid in reducing the incidence of seroma; and, in patients at high risk for post-operative infections, aid in reducing the incidence of superficial surgical site infection in Class I and Class II wounds.

The effectiveness of Prevena Therapy in reducing the incidence of SSIs and seroma in all surgical procedures and populations has not been demonstrated. See full indications for use and limitations at eifu.solventum.com.

The Solventum™ Prevena Restor™ Incision Management System is intended to manage the environment of surgical incisions that continue to drain following sutured or stapled closure by maintaining a closed environment and removing exudate via the application of negative pressure wound therapy.



Note: Specific indications, limitations, contraindications, warnings, precautions and safety information exist for these products and therapies. Please consult a clinician and product instructions for use prior to application. Rx only.

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