

Your closure is their beginning

Start your patient's recovery off right with a strong, protected closure from **DERMABOND® PRINEO® Skin Closure System**^{1,2}

Watertight seal³

Greater skin-holding strength than sutures or staples^{2,4*}



Microbial barrier protection⁵



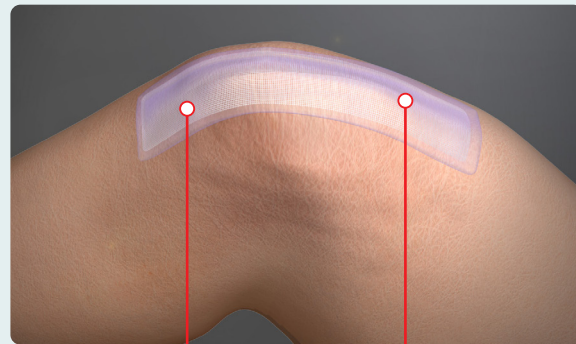
DERMABOND PRINEO Skin Closure System
is a non-invasive alternative to skin stitches and staples¹
 Help your patients focus on recovery

Protects incisions with a watertight, microbial barrier^{3,5}



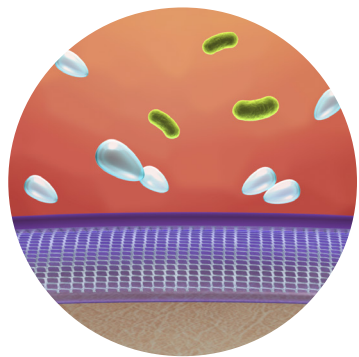
Provides a flexible microbial barrier with 99% protection in vitro for 72 hours against organisms commonly responsible for SSIs^{5,6†}

*Clinical significance unknown.



Self-adhering mesh

Liquid adhesive



Create an ideal wound-healing environment^{3,5,7}

- While **protecting the incision** against large elements like bacteria and water, DERMABOND PRINEO System allows smaller molecules, like water vapor, to pass through^{3,5,7}
- This **breathability** helps maintain a moist, wound-healing environment with an adequate amount of oxygen to further support wound healing⁷

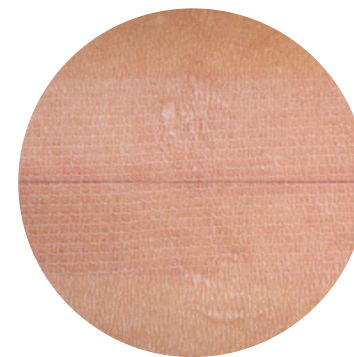
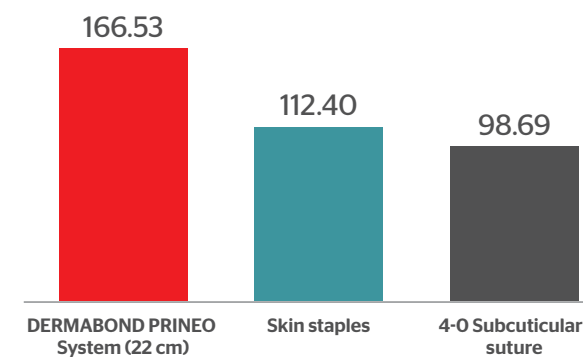
Give your patients the strength they need for optimal healing^{2,4}

Gain greater skin-holding strength than sutures or staples^{2,4§}

DERMABOND PRINEO System is **-33% and -40% stronger** compared to the average strength of staples and 4-0 suture, respectively^{2,4§}

COMPARATIVE SKIN-HOLDING STRENGTH^{2,4§}

Mean max load (N) prior to 3 mm gap (+/- 1 mm)



Provides even tension distribution—unlike sutures and staples

While sutures and staples penetrate the skin and place tension on the wounded tissue, DERMABOND PRINEO System **redistributes tension** across the surrounding healthy skin in a uniform way.

Not an actual patient; retouched photo for illustrative purposes only to show results post surgery. Individual result may vary.

Reduced rates of readmission⁸

In a retrospective study comparing DERMABOND PRINEO System and skin staples in total knee arthroplasty, DERMABOND PRINEO System was associated with **significantly reduced readmission rates** and probability of discharge to a skilled nursing facility or other non-home setting.^{8‡}

“[With DERMABOND PRINEO System], the less amount of wound contamination problems, [the less] patients have to come back to the clinic for wound checks.”

— **Dr. James E. Dowd**, Orthopaedic Surgeon, Virginia Beach, VA

The quote is the opinion of Dr. Dowd, a real surgeon who used DERMABOND PRINEO System. Dr. Dowd is a paid consultant of Ethicon. Post-surgical interview was May 8, 2017.

¹Challenge organisms included Staphylococcus epidermidis, Escherichia coli, Staphylococcus aureus, Pseudomonas aeruginosa, and Enterococcus faecium.
[†]Premier Inpatient Database in Total Knee Arthroplasty (p<0.05). N=1942; 2010-2015.

[§]In an ex-vivo study, more load in N was required to create a 3±1 mm gap between skin edges approximated with DERMABOND PRINEO System, than with subcuticular 4-0 MONOCRYL® Suture or PROXIMATE® staples (P=0.00).

Protection. Strength. Comfort. **All in one.**^{1,2,9*}

DERMABOND PRINEO Skin Closure System is the surgeon- and patient-preferred alternative to staples that combines proven microbial barrier protection and strength with a more comfortable recovery.^{1,2,9*}

- Delivers better cosmesis compared to staples^{9*}
- No bandages to change^{1,10}
- Patients can shower right away with your approval¹
- Leads to greater overall patient satisfaction when compared to staples^{9*}



For more information, ask your Ethicon representative or visit www.ethicon.com/optimalhealing.

For complete indications, contraindications, warnings, precautions, and adverse reactions, please reference full package insert.

*Double-blinded quantitative market research study comparing patient and surgeon experience with DERMABOND PRINEO System and skin staples in total knee arthroplasty. N=88 patients. N=83 orthopaedic surgeons. 90% c.i. Fielded June/July 2017.

References: 1. DERMABOND® PRINEO® Skin Closure System. Instructions for Use. Ethicon, Inc. 2. Kumar A. AST-2012-0290: Study to compare the tissue holding strength of PRINEO™ skin closure system with conventional wound closure techniques. October 11, 2012. Ethicon, Inc. 3. Kumar A. AST-2014-0060: Completion Report for Design Verification testing for DERMABOND™ PRINEO™ 22 cm skin closure system (DP22) AST-2014-0060, Version 2. April 19, 2016. Ethicon, Inc. 4. Kumar A. AST-2014-0246: Study to compare the tissue holding strengths of DERMABOND® PRINEO® 22 cm Skin Closure System (DP22) to conventional wound closure techniques. September 25, 2014. Ethicon, Inc. 5. Su W. Study Report for in vitro evaluation of microbial barrier properties of DERMABOND ProTape. Report Number O6TRO71. December 4, 2006. Ethicon, Inc. 6. Shapiro A, Dinsmore R, North J. Tensile strength of wound closure with cyanoacrylate glue. *Am Surg.* 2001;67:1113-1115. 7. Kannon GA, Garret AB. Moist wound healing with occlusive dressings. *Dermatol Surg.* 1995;21(7):583-590. 8. Sutton N, Schmitz ND, Johnston SS. Economic and clinical comparison of 2-octyl cyanoacrylate/polymer mesh tape with skin staples in total knee replacement. *J Wound Care.* 2018;27(Sup4):S12-S22. 9. PRINEO Claims Research Quant Detail. August 16, 2017. Ethicon, Inc. 10. De Cock E, van Nooten F, Mueller K, Tan R. Changing the surgical wound closure management pathway: time and supplies with PRINEO vs. standard of care for abdominoplasty surgery in Germany. Poster presented at: International Society for Pharmacoeconomics and Outcomes Research, 11th Annual European Congress: November 2008, Athens, Greece.