

K081580 BEAMER ARGON SNARE PROBEJun 19, 2008
14 days to decisionK081580 · Product code: **GEI** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k081580/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jun 5, 2008
Decision date	Jun 19, 2008
Days to decision	14 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	Conmed Corporation
Location	Utica, NY, US
Contact	KAREN PROVENCHER
Website	https://www.conmed.com
510(k) history	83 submissions · 83 cleared · 2004-2026

Conmed Corporation is a global medical device manufacturer specializing in surgical equipment and operating room solutions. The company operates with a manufacturing facility in Utica, US, and serves multiple surgical specialties including general surgery, orthopedics, and patient monitoring. Conmed has received FDA 510(k) clearances from total submissions since its first clearance in 2004. The company maintains active regulatory engagement, with its most recent clearance in 2026. Its cleared devices focus primarily on General & Plastic Surgery applications, including ele...
