FDA 510(k) Premarket Notification

K081644 BEAMER ARGON PROBE

CLEARED

Sep 10, 2008
90 days to decision

K081644 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k081644/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jun 12, 2008
Decision date	Sep 10, 2008
Days to decision	90 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...