

K103665 CONMED R SABRE GENESIS TM ELECTROSURGICAL UNITFeb 16, 2011
62 days to decisionK103665 · Product code: **GEI** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k103665/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Dec 16, 2010
Decision date	Feb 16, 2011
Days to decision	62 days
Third-party review	No
Summary / Statement	Summary
Other names	CONED R SABRE GENESIS TM ELECTROSURGICAL UNIT (230V); CONMED HYFRESURG O

APPLICANT

Company	Conmed Corporation
Location	Utica, NY, US
Contact	SHAWN RIEDEL
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...

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k103665/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 19, 2026