

K052009 CONMED SYSTEM MODEL 2450 ELECTROSURGICAL GENERATORSep 28, 2005
65 days to decisionK052009 · Product code: **GEI** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k052009/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jul 25, 2005
Decision date	Sep 28, 2005
Days to decision	65 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Conmed Corporation
Location	Utica, NY, US
Contact	PAMELA VETTER
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...

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Device record: <https://www.510kdatabase.net/k052009/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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