

**K081634 CONMED GOLDVAC INTEGRATED SMOKE  
EVACUATION PENCIL**Jun 25, 2008  
14 days to decisionK081634 · Product code: **GEI** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k081634/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jun 11, 2008
Decision date	Jun 25, 2008
Days to decision	14 days
Third-party review	Yes
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Conmed Corporation</b>
Location	Utica, NY, US
Contact	SHAWN RIEDEL
Website	<a href="https://www.conmed.com">https://www.conmed.com</a>
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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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k081634/>. Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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