

**K083781 EEA HEMORRHOID STAPLER & ACCESSORIES**Feb 4, 2009  
47 days to decisionK083781 · Product code: **GDW** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k083781/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Staple, Implantable (GDW)
Date received	Dec 19, 2008
Decision date	Feb 4, 2009
Days to decision	47 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Surgical Devices, A Global Business Unit Covidien</b>
Location	North Haven, CT, US
Contact	FRANK GIANELLI
510(k) history	3 submissions · 3 cleared · 2008-2009

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k083781/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 23, 2026