

**K060255 E SURGICAL ELECTROSURGICAL PATIENT RETURN ELECTRODE, DUAL PLATE WITH CORD AND ELECTROSURGICAL PATIENT RETURN ELECTRODE**

Mar 31, 2006  
58 days to decision

K060255 · Product code: **GEI** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k060255/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Feb 1, 2006
Decision date	Mar 31, 2006
Days to decision	58 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>E Surgical, LLC</b>
Location	Boulder, CO, US
Contact	LEWIS WARD
510(k) history	5 submissions · 5 cleared · 2006-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k060255/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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