

**K823083 ELECTROSURGICAL UNIT**Dec 7, 1982  
49 days to decisionK823083 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k823083/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Oct 19, 1982
Decision date	Dec 7, 1982
Days to decision	49 days
Third-party review	No

**APPLICANT**

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Company	<b>Kelleher Corp.</b>
Location	Mchenry, IL, US
510(k) history	94 submissions · 94 cleared · 1982-1983

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Device record: <https://www.510kdatabase.net/k823083/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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