

**K771639 ELECTROSURGICAL GENERATOR**Nov 29, 1977  
95 days to decisionK771639 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k771639/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Aug 26, 1977
Decision date	Nov 29, 1977
Days to decision	95 days
Third-party review	No

**APPLICANT**

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Company	<b>Cameron-Miller, Inc.</b>
Location	Mchenry, IL, US
510(k) history	14 submissions · 14 cleared · 1977-1994

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

Device record: <https://www.510kdatabase.net/k771639/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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