

**K812702 TECHNICATOR**Nov 5, 1981  
43 days to decisionK812702 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k812702/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Sep 23, 1981
Decision date	Nov 5, 1981
Days to decision	43 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/k812702/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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