

**K001382 AARON 2100 HIGH FREQUENCY ELECTROSURGICAL GENERATOR**Jun 27, 2000  
56 days to decisionK001382 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k001382/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	May 2, 2000
Decision date	Jun 27, 2000
Days to decision	56 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Aaron Medical Industries</b>
Location	St. Petersburg, FL, US
Contact	RICK KOZLOFF
510(k) history	46 submissions · 46 cleared · 1990-2006

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k001382/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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