

**K070101 UNIBLATE ELECTROSURGICAL DEVICE**Feb 14, 2007  
35 days to decisionK070101 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k070101/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jan 10, 2007
Decision date	Feb 14, 2007
Days to decision	35 days
Third-party review	Yes
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Rita Medical Systems, Inc.</b>
Location	Mountain View, CA, US
Contact	KAM H LEUNG
510(k) history	9 submissions · 9 cleared · 2003-2007

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k070101/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 28, 2026