

**K030049 MODIFICATION TO RUBICOR ENCAPSULE BREAST BIOPSY DEVICE, MODEL 30086**Jan 29, 2003  
23 days to decisionK030049 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k030049/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jan 6, 2003
Decision date	Jan 29, 2003
Days to decision	23 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Rubicor Medical, Inc.</b>
Location	San Carlos, CA, US
Contact	ROBERT J CHIN
510(k) history	8 submissions · 8 cleared · 2002-2007

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k030049/>; 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