

**K071048 RUBICOR MAGIC BREAST BIOPSY DEVICE, MODEL  
31537**May 15, 2007  
32 days to decisionK071048 · Product code: **KNW** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k071048/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Instrument, Biopsy (KNW)
Date received	Apr 13, 2007
Decision date	May 15, 2007
Days to decision	32 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/k071048/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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