FDA 510(k) Premarket Notification

# K071048 RUBICOR MAGIC BREAST BIOPSY DEVICE, MODEL 31537

**CLEARED**

May 15, 2007
32 days to decision

K071048 · 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

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