FDA 510(k) Premarket Notification

K030049 MODIFICATION TO RUBICOR ENCAPSULE BREAST BIOPSY DEVICE, MODEL 30086

CLEARED

Jan 29, 2003
23 days to decision

K030049 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k030049/

SUBMISSION DETAILS

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

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