FDA 510(k) Premarket Notification

K021329 RITA SYSTEM

CLEARED

Oct 9, 2002
166 days to decision

K021329 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k021329/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Apr 26, 2002
Decision date	Oct 9, 2002
Days to decision	166 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Rita Medical Systems**
Location	Mountain View, CA, US
Contact	ERIN MAZZONE
510(k) history	11 submissions · 10 cleared · 1998-2006

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k021329/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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