FDA 510(k) Premarket Notification

K013904 BMC RADIO FREQUENCY PERFORATION GENERATOR, MODELS RFP-100-115 AND RFP-100-230

CLEARED

Feb 12, 2002
78 days to decision

K013904 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k013904/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Nov 26, 2001
Decision date	Feb 12, 2002
Days to decision	78 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Baylis Medical Co., Inc.**
Location	Mississauga, CA
Contact	KRIS SHAH
510(k) history	28 submissions · 28 cleared · 1998-2013

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k013904/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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