FDA 510(k) Premarket Notification

K023288 COBRA BIPOLAR CLAMP, MODEL 15902

CLEARED

Apr 2, 2003
182 days to decision

K023288 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k023288/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Oct 2, 2002
Decision date	Apr 2, 2003
Days to decision	182 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Boston Scientific EP Technologies**
Location	San Jose, CA, US
Contact	APRIL MALMBORG
510(k) history	1 submissions · 1 cleared · 2003-2003

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k023288/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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