FDA 510(k) Premarket Notification

K012018 BICAP COAG BIPOLAR LAPAROSCOPY PROBE, MODEL 006908-910

CLEARED

Jul 16, 2002
383 days to decision

K012018 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k012018/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jun 28, 2001
Decision date	Jul 16, 2002
Days to decision	383 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Acmi Corporation**
Location	Southborough, MA, US
Contact	FRANK J FUCILE
510(k) history	16 submissions · 16 cleared · 2002-2006