

**K012018 BICAP COAG BIPOLAR LAPAROSCOPY PROBE,
MODEL 006908-910**Jul 16, 2002
383 days to decisionK012018 · Product code: **GEI** · General & Plastic SurgerySource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k012018/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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