

K973190 ACCUPROBE 450, ACCUPROBE 550/530 AND THE ACCUPROBE 600 SERIES

Nov 21, 1997
88 days to decision

K973190 · Product code: **GEI** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k973190/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Aug 25, 1997
Decision date	Nov 21, 1997
Days to decision	88 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Cryomedical Sciences, Inc.
Location	Bethesda, MD, US
Contact	RICHARD J REINHART
510(k) history	10 submissions · 10 cleared · 1991-2001

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Device record: <https://www.510kdatabase.net/k973190/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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