FDA 510(k) Premarket Notification

K972350 MODEL 50 BIPOLAR COAGULATOR

CLEARED

Aug 21, 1997
58 days to decision

K972350 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k972350/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jun 24, 1997
Decision date	Aug 21, 1997
Days to decision	58 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Boston Surgical Products, Inc.**
Location	Kingston, MA, US
Contact	CHARLES VASSALLO
510(k) history	17 submissions · 17 cleared · 1994-1997