FDA 510(k) Premarket Notification

K801153 VALLEYLAB RETURN ELECTRODE MONITOR

CLEARED

May 20, 1980
6 days to decision

K801153 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k801153/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	May 14, 1980
Decision date	May 20, 1980
Days to decision	6 days
Third-party review	No

APPLICANT

Company	**Valleylab, Inc.**
Location	Mchenry, IL, US
510(k) history	94 submissions · 93 cleared · 1976-2003