FDA 510(k) Premarket Notification

K953737 VL7600 REM PATIENT RETURN ELECTRODE

CLEARED

Aug 23, 1995
13 days to decision

K953737 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k953737/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Aug 10, 1995
Decision date	Aug 23, 1995
Days to decision	13 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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