FDA 510(k) Premarket Notification

K813072 E2502 & E2502B LECTROSWITCH

CLEARED

Nov 10, 1981
8 days to decision

K813072 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k813072/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Nov 2, 1981
Decision date	Nov 10, 1981
Days to decision	8 days
Third-party review	No

APPLICANT

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