FDA 510(k) Premarket Notification

K822572 POLYHESIVE PATIENT RETURN ELECTRODE

CLEARED

Sep 21, 1982
28 days to decision

K822572 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k822572/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Aug 24, 1982
Decision date	Sep 21, 1982
Days to decision	28 days
Third-party review	No

APPLICANT

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