FDA 510(k) Premarket Notification

K964602 DAVIS BAYONET ELECTRODES

CLEARED

Jan 9, 1997
52 days to decision

K964602 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k964602/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Nov 18, 1996
Decision date	Jan 9, 1997
Days to decision	52 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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