FDA 510(k) Premarket Notification

K822513 MODEL #2000 HI-TEMP & #3000 LOW-TEMP

CLEARED

Sep 21, 1982
32 days to decision

K822513 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k822513/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Aug 20, 1982
Decision date	Sep 21, 1982
Days to decision	32 days
Third-party review	No

APPLICANT

Company	**Surgicare Scientific Laboratory, Inc.**
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1982-1983