FDA 510(k) Premarket Notification

K823083 ELECTROSURGICAL UNIT

CLEARED

Dec 7, 1982
49 days to decision

K823083 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k823083/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Oct 19, 1982
Decision date	Dec 7, 1982
Days to decision	49 days
Third-party review	No

APPLICANT

Company	**Kelleher Corp.**
Location	Mchenry, IL, US
510(k) history	94 submissions · 94 cleared · 1982-1983