FDA 510(k) Premarket Notification

K792600 MODEL III HANDSWITCH

CLEARED

Jan 10, 1980
23 days to decision

K792600 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k792600/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Dec 18, 1979
Decision date	Jan 10, 1980
Days to decision	23 days
Third-party review	No

APPLICANT

Company	**Sara Dan, Inc.**
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1979-1980