FDA 510(k) Premarket Notification

K812702 TECHNICATOR

CLEARED

Nov 5, 1981
43 days to decision

K812702 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k812702/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Sep 23, 1981
Decision date	Nov 5, 1981
Days to decision	43 days
Third-party review	No

APPLICANT

Company	**Cameron-Miller, Inc.**
Location	Mchenry, IL, US
510(k) history	14 submissions · 14 cleared · 1977-1994