FDA 510(k) Premarket Notification

K820902 MULTIPERFECTEMP

CLEARED

Apr 26, 1982
26 days to decision

K820902 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k820902/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Mar 31, 1982
Decision date	Apr 26, 1982
Days to decision	26 days
Third-party review	No

APPLICANT

Company	**Kay Medical, Inc.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1982-1982