FDA 510(k) Premarket Notification

K802942 HITEMP. TM 2 LOOPTIP CAUTERY

CLEARED

Feb 4, 1981
77 days to decision

K802942 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k802942/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Nov 19, 1980
Decision date	Feb 4, 1981
Days to decision	77 days
Third-party review	No

APPLICANT

Company	**Medical Products Development, Inc.**
Location	Mchenry, IL, US
510(k) history	13 submissions · 12 cleared · 1980-1981