FDA 510(k) Premarket Notification

K932817 HM-880 I

CLEARED

Mar 9, 1994
273 days to decision

K932817 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k932817/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jun 9, 1993
Decision date	Mar 9, 1994
Days to decision	273 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Hill-Med, Inc.**
Location	Miami, FL, US
Contact	ERNESTO ACKERMAN
510(k) history	10 submissions · 10 cleared · 1993-2000