FDA 510(k) Premarket Notification

K890234 SUTURE CLIP REMOVER

CLEARED

Jan 27, 1989
9 days to decision

K890234 · Product code: **HTD** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k890234/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Forceps (HTD)
Date received	Jan 18, 1989
Decision date	Jan 27, 1989
Days to decision	9 days
Third-party review	No

APPLICANT

Company	**Kinetic Medical Products**
Location	Erie, PA, US
Contact	JAMES I LAUGHNER
510(k) history	63 submissions · 63 cleared · 1989-1989