FDA 510(k) Premarket Notification

K854890 HOOK PROBE

CLEARED

May 2, 1986
147 days to decision

K854890 · Product code: **KDC** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k854890/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Surgical, Disposable (KDC)
Date received	Dec 6, 1985
Decision date	May 2, 1986
Days to decision	147 days
Third-party review	No

APPLICANT

Company	**Euro-Med Intl.**
Location	Danville, CA, US
Contact	DONALD HOLSTEN
510(k) history	16 submissions · 16 cleared · 1985-1988