FDA 510(k) Premarket Notification

K881276 HR-5000 ELECTROSURGICAL DEVICE FOR NEW INDICATIONS

CLEARED

May 4, 1988
40 days to decision

K881276 · Product code: **KCW** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k881276/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Epilator, High Frequency, Needle-type (KCW)
Date received	Mar 25, 1988
Decision date	May 4, 1988
Days to decision	40 days
Third-party review	No

APPLICANT

Company	**Ime Co. , Ltd.**
Location	Washington, DC, US
Contact	KEVIN G MCANANEY
510(k) history	2 submissions · 2 cleared · 1987-1988