FDA 510(k) Premarket Notification

K874138 ACCUCOM 2

CLEARED

Jun 1, 1988
232 days to decision

K874138 · Product code: **DPT** · Cardiovascular
Source: https://www.510kdatabase.net/k874138/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Probe, Blood-flow, Extravascular (DPT)
Date received	Oct 13, 1987
Decision date	Jun 1, 1988
Days to decision	232 days
Third-party review	No

APPLICANT

Company	**Datascope Corp.**
Location	Mchenry, IL, US
Contact	ARTHUR FRIEDMAN
510(k) history	136 submissions · 135 cleared · 1976-2019