FDA 510(k) Premarket Notification

K881455 EMBOLECTOMY CATHETER(MODIFIED)

CLEARED

Jun 15, 1988
70 days to decision

K881455 · Product code: **DXE** · Cardiovascular
Source: https://www.510kdatabase.net/k881455/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Embolectomy (DXE)
Date received	Apr 6, 1988
Decision date	Jun 15, 1988
Days to decision	70 days
Third-party review	No

APPLICANT

Company	**Cathlab Corp.**
Location	Tustin, CA, US
Contact	SATTAR SAUDAGAR
510(k) history	8 submissions · 8 cleared · 1988-1991