FDA 510(k) Premarket Notification

K832850 ELECATH PACEWEDGE DUAL PRESS. BIPOLAR

CLEARED

Oct 19, 1983
58 days to decision

K832850 · Product code: **LDF** · Cardiovascular
Source: https://www.510kdatabase.net/k832850/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	Aug 22, 1983
Decision date	Oct 19, 1983
Days to decision	58 days
Third-party review	No

APPLICANT

Company	**Electro-Catheter Corp.**
Location	Mchenry, IL, US
510(k) history	35 submissions · 35 cleared · 1976-1995