FDA 510(k) Premarket Notification

K810312 ELECATH DUAL MODE TRANSTHORACIC PACING

CLEARED

Mar 4, 1981
27 days to decision

K810312 · Product code: **LDF** · Cardiovascular
Source: https://www.510kdatabase.net/k810312/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	Feb 5, 1981
Decision date	Mar 4, 1981
Days to decision	27 days
Third-party review	No

APPLICANT

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