

K832850 ELECATH PACEWEDGE DUAL PRESS. BIPOLAROct 19, 1983
58 days to decisionK832850 · Product code: **LDF** · CardiovascularSource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k832850/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 8, 2026