

K810312 ELECATH DUAL MODE TRANSTHORACIC PACINGMar 4, 1981
27 days to decisionK810312 · Product code: **LDF** · CardiovascularSource: <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

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Device record: <https://www.510kdatabase.net/k810312/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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