

K812517 CARDIAC PACING ELECTRODESOct 13, 1981
41 days to decisionK812517 · Product code: **LDF** · CardiovascularSource: <https://www.510kdatabase.net/k812517/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	Sep 2, 1981
Decision date	Oct 13, 1981
Days to decision	41 days
Third-party review	No

APPLICANT

Company	Medics Instrument Corp.
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1981-1991

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Device record: <https://www.510kdatabase.net/k812517/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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