

K900982 E-CLIPApr 19, 1990
48 days to decisionK900982 · Product code: **LDF** · Cardiovascular
Source: <https://www.510kdatabase.net/k900982/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	Mar 2, 1990
Decision date	Apr 19, 1990
Days to decision	48 days
Third-party review	No

APPLICANT

Company	Joel Cohen & Esther R. Cohen
Location	Plantation, FL, US
Contact	JOEL COHEN
510(k) history	1 submissions · 1 cleared · 1990-1990

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k900982/>; 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 6, 2026