

K890018 PACEMAKER LEAD ADAPTERJun 20, 1989
168 days to decisionK890018 · Product code: **DTB** · CardiovascularSource: <https://www.510kdatabase.net/k890018/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Permanent Pacemaker Electrode (DTB)
Date received	Jan 3, 1989
Decision date	Jun 20, 1989
Days to decision	168 days
Third-party review	No

APPLICANT

Company	Ela Medical, Inc.
Location	Mchenry, IL, US
Contact	PAUL KOHL
510(k) history	43 submissions · 36 cleared · 1979-2004

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k890018/>; 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 3, 2026