

**K896742 FOCUS J81F, J83F, J41F, J43F PACING LEADS**Jul 17, 1990  
229 days to decisionK896742 · Product code: **DTB** · CardiovascularSource: <https://www.510kdatabase.net/k896742/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Permanent Pacemaker Electrode (DTB)
Date received	Nov 30, 1989
Decision date	Jul 17, 1990
Days to decision	229 days
Third-party review	No

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k896742/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated July 3, 2026