

**K884775 ARBO PINK H82P**Feb 6, 1989  
82 days to decisionK884775 · Product code: **DRX** · CardiovascularSource: <https://www.510kdatabase.net/k884775/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Nov 16, 1988
Decision date	Feb 6, 1989
Days to decision	82 days
Third-party review	No

**APPLICANT**

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Company	<b>Arbo Medical, Inc.</b>
Location	Wilton, CT, US
Contact	WAYNE D SHOCKLOSS
510(k) history	15 submissions · 15 cleared · 1989-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k884775/>, 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 2, 2026