

**K935246 ARBO H49P, H59P DISPOSABLE MONITORING ELECTRODES**Nov 16, 1994  
380 days to decisionK935246 · Product code: **DRX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k935246/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Nov 1, 1993
Decision date	Nov 16, 1994
Days to decision	380 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Arbo Medical, Inc.</b>
Location	Wilton, CT, US
Contact	JEFFREY M BLOCK
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/k935246/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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