

**K860298 STABLELINE**Apr 2, 1986  
64 days to decisionK860298 · Product code: **KRI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k860298/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accessory Equipment, Cardiopulmonary Bypass (KRI)
Date received	Jan 28, 1986
Decision date	Apr 2, 1986
Days to decision	64 days
Third-party review	No

**APPLICANT**

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Company	<b>Cardio Metrics, Inc.</b>
Location	Houston, TX, US
Contact	MARY K NOONEN
510(k) history	23 submissions · 23 cleared · 1986-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k860298/>; 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 May 20, 2026