

**K821090 CARDI-O-LIFE MODEL 8823**Jul 8, 1982  
80 days to decisionK821090 · Product code: **DPS** · CardiovascularSource: <https://www.510kdatabase.net/k821090/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrocardiograph (DPS)
Date received	Apr 19, 1982
Decision date	Jul 8, 1982
Days to decision	80 days
Third-party review	No

**APPLICANT**

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Company	<b>Amertek</b>
Location	Walker, MI, US
510(k) history	2 submissions · 2 cleared · 1981-1982

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k821090/> 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 June 29, 2026