

**K852181 ELECTROCARDIOGRAPH**Oct 9, 1985  
142 days to decisionK852181 · Product code: **DPS** · Cardiovascular  
Source: <https://www.510kdatabase.net/k852181/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrocardiograph (DPS)
Date received	May 20, 1985
Decision date	Oct 9, 1985
Days to decision	142 days
Third-party review	No

**APPLICANT**

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Company	<b>Armstrong Ind., Inc.</b>
Location	Mchenry, IL, US
Contact	KAREN BOWERS
510(k) history	5 submissions · 5 cleared · 1982-1985

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k852181/>; 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 27, 2026