

**K782033 CARDIAC CARE Q/C TEST KIT**Dec 15, 1978  
9 days to decisionK782033 · Product code: **DRL** · CardiovascularSource: <https://www.510kdatabase.net/k782033/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tester, Defibrillator (DRL)
Date received	Dec 6, 1978
Decision date	Dec 15, 1978
Days to decision	9 days
Third-party review	No

**APPLICANT**

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Company	<b>Instrumentation Laboratory CO</b>
Location	Mchenry, IL, US
510(k) history	321 submissions · 320 cleared · 1976-2023

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k782033/> 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 15, 2026