

**K820579 FLEX-E1 ECG ELECTRODE**Apr 14, 1982  
42 days to decisionK820579 · Product code: **DRS** · CardiovascularSource: <https://www.510kdatabase.net/k820579/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Blood-pressure, Extravascular (DRS)
Date received	Mar 3, 1982
Decision date	Apr 14, 1982
Days to decision	42 days
Third-party review	No

**APPLICANT**

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Company	<b>Conmedcorp</b>
Location	Dayton, OH, US
510(k) history	92 submissions · 92 cleared · 1981-2010

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k820579/> 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