

**K813358 GERARD MEDICAL ELECTRODE TESTER**Dec 29, 1981  
29 days to decisionK813358 · Product code: **KRC** · CardiovascularSource: <https://www.510kdatabase.net/k813358/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tester, Electrode, Surface, Electrocardiographic (KRC)
Date received	Nov 30, 1981
Decision date	Dec 29, 1981
Days to decision	29 days
Third-party review	No

**APPLICANT**

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Company	<b>Gerard Medical Enterprises, Inc.</b>
Location	Mchenry, IL, US
510(k) history	8 submissions · 6 cleared · 1981-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k813358/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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