

K780501 ELECTRODE TESTERMay 19, 1978
52 days to decisionK780501 · Product code: **KRC** · CardiovascularSource: <https://www.510kdatabase.net/k780501/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tester, Electrode, Surface, Electrocardiographic (KRC)
Date received	Mar 28, 1978
Decision date	May 19, 1978
Days to decision	52 days
Third-party review	No

APPLICANT

Company	Mg Medical Electronics
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1978-1978

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Device record: <https://www.510kdatabase.net/k780501/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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