

K833619 CONDUCTANCE METERJan 27, 1984
105 days to decisionK833619 · Product code: **KRC** · CardiovascularSource: <https://www.510kdatabase.net/k833619/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tester, Electrode, Surface, Electrocardiographic (KRC)
Date received	Oct 14, 1983
Decision date	Jan 27, 1984
Days to decision	105 days
Third-party review	No

APPLICANT

Company	Circadian, Inc.
Location	Mchenry, IL, US
510(k) history	13 submissions · 13 cleared · 1980-1992

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Device record: <https://www.510kdatabase.net/k833619/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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