

K802182 WISCONSIN HEART RATE MONITOROct 31, 1980
52 days to decisionK802182 · Product code: **DRT** · CardiovascularSource: <https://www.510kdatabase.net/k802182/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiometer & Rate Alarm) (DRT)
Date received	Sep 9, 1980
Decision date	Oct 31, 1980
Days to decision	52 days
Third-party review	No

APPLICANT

Company	Lunar Radiation Corp.
Location	WI, US
510(k) history	6 submissions · 6 cleared · 1980-1988

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Device record: <https://www.510kdatabase.net/k802182/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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