

K830830 CIRCA MED TWO-CHANNELJun 18, 1983
94 days to decisionK830830 · Product code: **DSI** · CardiovascularSource: <https://www.510kdatabase.net/k830830/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Mar 16, 1983
Decision date	Jun 18, 1983
Days to decision	94 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/k830830/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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