

K822718 SERIES 7000 BEDSIDE PATIENT MONITOROct 4, 1982
27 days to decisionK822718 · Product code: **DSI** · CardiovascularSource: <https://www.510kdatabase.net/k822718/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Sep 7, 1982
Decision date	Oct 4, 1982
Days to decision	27 days
Third-party review	No

APPLICANT

Company	Marquette Electronics, Inc.
Location	Walker, MI, US
510(k) history	82 submissions · 81 cleared · 1980-1997

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Device record: <https://www.510kdatabase.net/k822718/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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