

K822030 MULTIPLE LEAD ARRHYTHMIA MODULESep 28, 1982
78 days to decisionK822030 · Product code: **DSI** · Cardiovascular
Source: <https://www.510kdatabase.net/k822030/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Jul 12, 1982
Decision date	Sep 28, 1982
Days to decision	78 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k822030/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 21, 2026