

K954852 SOLAR 9000 ANESTHESIA INFORMATION MONITORMay 29, 1996
219 days to decisionK954852 · Product code: **DSI** · Cardiovascular
Source: <https://www.510kdatabase.net/k954852/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Oct 23, 1995
Decision date	May 29, 1996
Days to decision	219 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k954852/> 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