FDA 510(k) Premarket Notification

K822030 MULTIPLE LEAD ARRHYTHMIA MODULE

CLEARED

Sep 28, 1982
78 days to decision

K822030 · 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