FDA 510(k) Premarket Notification

K822718 SERIES 7000 BEDSIDE PATIENT MONITOR

CLEARED

Oct 4, 1982
27 days to decision

K822718 · Product code: **DSI** · Cardiovascular
Source: 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