FDA 510(k) Premarket Notification

K872354 SERIES 2000 BEDSIDE PATIENT MONITOR

CLEARED

Jul 21, 1987
33 days to decision

K872354 · Product code: **DRT** · Cardiovascular
Source: https://www.510kdatabase.net/k872354/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Jun 18, 1987
Decision date	Jul 21, 1987
Days to decision	33 days
Third-party review	No

APPLICANT

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