FDA 510(k) Premarket Notification

K780432 ARRHYTHMIA PROGRAM MODULE

CLEARED

May 26, 1978
70 days to decision

K780432 · Product code: **DSI** · Cardiovascular
Source: https://www.510kdatabase.net/k780432/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Mar 17, 1978
Decision date	May 26, 1978
Days to decision	70 days
Third-party review	No

APPLICANT

Company	**Midwest Analog & Digital, Inc.**
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1978-1985