FDA 510(k) Premarket Notification

K772360 ARRHYTHMIA MODULE

CLEARED

Jan 12, 1978
16 days to decision

K772360 · Product code: **DSI** · Cardiovascular
Source: https://www.510kdatabase.net/k772360/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Dec 27, 1977
Decision date	Jan 12, 1978
Days to decision	16 days
Third-party review	No

APPLICANT

Company	**General Electric Co.**
Location	Mchenry, IL, US
510(k) history	254 submissions · 254 cleared · 1976-2011