FDA 510(k) Premarket Notification

K821173 EVENTMASTER 3 HOLTER EVENT SCANNER

CLEARED

May 18, 1982
22 days to decision

K821173 · Product code: **DSF** · Cardiovascular
Source: https://www.510kdatabase.net/k821173/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Recorder, Paper Chart (DSF)
Date received	Apr 26, 1982
Decision date	May 18, 1982
Days to decision	22 days
Third-party review	No

APPLICANT

Company	**Instruments For Cardiac Research**
Location	Mchenry, IL, US
510(k) history	8 submissions · 8 cleared · 1977-1984