FDA 510(k) Premarket Notification

K833162 AMBULATORY MONITORING DEVICE

CLEARED

Jan 10, 1984
116 days to decision

K833162 · Product code: **DSI** · Cardiovascular
Source: https://www.510kdatabase.net/k833162/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Sep 16, 1983
Decision date	Jan 10, 1984
Days to decision	116 days
Third-party review	No

APPLICANT

Company	**Oxford Medilog, Inc.**
Location	Mchenry, IL, US
510(k) history	48 submissions · 48 cleared · 1978-1994