FDA 510(k) Premarket Notification

K802033 OXFORD INTENSIVE CARE PATIENT MON. SYS

CLEARED

Sep 26, 1980
35 days to decision

K802033 · Product code: **DSI** · Cardiovascular
Source: https://www.510kdatabase.net/k802033/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Aug 22, 1980
Decision date	Sep 26, 1980
Days to decision	35 days
Third-party review	No

APPLICANT

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