FDA 510(k) Premarket Notification

K790287 ANALYZER, MEDILOG II

CLEARED

Feb 28, 1979
20 days to decision

K790287 · Product code: **DSI** · Cardiovascular
Source: https://www.510kdatabase.net/k790287/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Feb 8, 1979
Decision date	Feb 28, 1979
Days to decision	20 days
Third-party review	No

APPLICANT

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