FDA 510(k) Premarket Notification

K992323 DATEX-OHMEDA CARDIOCAP 5, F-MX, F-MXG AND ACCESSORIES

CLEARED

Nov 1, 1999
112 days to decision

K992323 · Product code: **MLD** · Cardiovascular
Source: https://www.510kdatabase.net/k992323/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, St Segment With Alarm (MLD)
Date received	Jul 12, 1999
Decision date	Nov 1, 1999
Days to decision	112 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Datex-Ohmeda, Inc.**
Location	Madison, WI, US
Contact	JOEL C KENT
510(k) history	60 submissions · 60 cleared · 1998-2025