

K992323 DATEX-OHMEDA CARDIOCAP 5, F-MX, F-MXG AND ACCESSORIESNov 1, 1999
112 days to decisionK992323 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k992323/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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