

K014255 DINAMAP PROCARE SERIES MONITORJul 2, 2002
188 days to decisionK014255 · Product code: **MWI** · Cardiovascular
Source: <https://www.510kdatabase.net/k014255/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient (without Arrhythmia Detection Or Alarms) (MWI)
Date received	Dec 26, 2001
Decision date	Jul 2, 2002
Days to decision	188 days
Third-party review	No
Summary / Statement	Summary
Other names	MODEL 100, 200, 300, 400

APPLICANT

Company	Ge Medical Systems Information Technologies
Location	Mchenry, IL, US
Contact	THOMAS ENGLISH
510(k) history	136 submissions · 132 cleared · 1978-2012

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k014255/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 26, 2026