

**K910665 MODEL 556 MONITOR W/24-HR. TREND
RECALL/MODIFIED**Dec 31, 1991
323 days to decisionK910665 · Product code: **DRT** · Cardiovascular
Source: <https://www.510kdatabase.net/k910665/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Feb 11, 1991
Decision date	Dec 31, 1991
Days to decision	323 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k910665/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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