

**K001112 GE MARQUETTE CLINICAL INFORMATION CENTER,
GE MARQUETTE CIC**Jun 14, 2000
69 days to decisionK001112 · Product code: **DXJ** · Cardiovascular
Source: <https://www.510kdatabase.net/k001112/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Display, Cathode-ray Tube, Medical (DXJ)
Date received	Apr 6, 2000
Decision date	Jun 14, 2000
Days to decision	69 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	General Electric Medical Systems Information Techn
Location	Sugarland, TX, US
Contact	DAVID WAHLIG
510(k) history	33 submissions · 33 cleared · 1999-2002

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k001112/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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