FDA 510(k) Premarket Notification

CLEARED

K001112 GE MARQUETTE CLINICAL INFORMATION CENTER, GE MARQUETTE CIC

Jun 14, 2000
69 days to decision

K001112 · 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