FDA 510(k) Premarket Notification

K863153 MODIFICATION OF CARDIOINTEGRAM/CARDIOINTEGRAPH

CLEARED

Jan 6, 1987
144 days to decision

K863153 · Product code: **DXG** · Cardiovascular
Source: https://www.510kdatabase.net/k863153/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Computer, Diagnostic, Pre-programmed, Single-function (DXG)
Date received	Aug 15, 1986
Decision date	Jan 6, 1987
Days to decision	144 days
Third-party review	No

APPLICANT

Company	**Ocg Technology, Inc.**
Location	Mchenry, IL, US
Contact	STEPHEN RAKOCZY
510(k) history	3 submissions · 3 cleared · 1981-1987