

**K863153 MODIFICATION OF
CARDIOINTEGRAM/CARDIOINTEGRAPH**Jan 6, 1987
144 days to decisionK863153 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k863153/> 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 July 7, 2026