

K925844 PHILIPS SUPER CP FAMILYDec 28, 1992
40 days to decisionK925844 · Product code: **IZO** · Radiology
Source: <https://www.510kdatabase.net/k925844/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Generator, High-voltage, X-ray, Diagnostic (IZO)
Date received	Nov 18, 1992
Decision date	Dec 28, 1992
Days to decision	40 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Philips Medical Systems, Inc.
Location	Mchenry, IL, US
Contact	WILLIAM G MCMAHON
510(k) history	111 submissions · 110 cleared · 1983-2010

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k925844/> 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 May 17, 2026