FDA 510(k) Premarket Notification

K925844 PHILIPS SUPER CP FAMILY

CLEARED

Dec 28, 1992
40 days to decision

K925844 · 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