

K953910 PHILIPS BV 300 SERIESOct 18, 1995
58 days to decisionK953910 · Product code: **OXO** · Radiology
Source: <https://www.510kdatabase.net/k953910/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Image-intensified Fluoroscopic X-ray System, Mobile (OXO)
Date received	Aug 21, 1995
Decision date	Oct 18, 1995
Days to decision	58 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Philips Medical Systems North America, Inc.
Location	Shelton, CT, US
Contact	PETER ALTMAN
510(k) history	71 submissions · 71 cleared · 1989-2010

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k953910/> 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