FDA 510(k) Premarket Notification

K953910 PHILIPS BV 300 SERIES

CLEARED

Oct 18, 1995
58 days to decision

K953910 · 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