FDA 510(k) Premarket Notification

K945278 PHILIPS BUCKY DIAGNOST FAMILY

CLEARED

Dec 14, 1994
47 days to decision

K945278 · Product code: **KPR** · Radiology
Source: https://www.510kdatabase.net/k945278/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Stationary (KPR)
Date received	Oct 28, 1994
Decision date	Dec 14, 1994
Days to decision	47 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