FDA 510(k) Premarket Notification

K941931 DIAGNOSTIC X-RAY PATIENT EXPOSURE INDICATOR

CLEARED

May 26, 1994
36 days to decision

K941931 · Product code: **KPR** · Radiology
Source: https://www.510kdatabase.net/k941931/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Stationary (KPR)
Date received	Apr 20, 1994
Decision date	May 26, 1994
Days to decision	36 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Victoreen, Inc.**
Location	Mchenry, IL, US
Contact	LINDA S MORIN
510(k) history	40 submissions · 40 cleared · 1979-1998