FDA 510(k) Premarket Notification

K923455 HFG 350, 650, 1050

CLEARED

Dec 28, 1992
167 days to decision

K923455 · Product code: **KPR** · Radiology
Source: https://www.510kdatabase.net/k923455/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Stationary (KPR)
Date received	Jul 14, 1992
Decision date	Dec 28, 1992
Days to decision	167 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Varian Canada, Inc.**
Location	Canada L7g 2j4, CA
Contact	ANTHONY B BAILEY
510(k) history	7 submissions · 7 cleared · 1991-1995