FDA 510(k) Premarket Notification

K860469 37-701-1 PATIENT DOASE MONITOR WITH DETECTORS

CLEARED

Apr 16, 1986
69 days to decision

K860469 · Product code: **KPZ** · Radiology
Source: https://www.510kdatabase.net/k860469/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Generator, High Voltage, X-ray, Therapeutic (KPZ)
Date received	Feb 6, 1986
Decision date	Apr 16, 1986
Days to decision	69 days
Third-party review	No

APPLICANT

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