

K860469 37-701-1 PATIENT DOASE MONITOR WITH DETECTORSApr 16, 1986
69 days to decisionK860469 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k860469/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 19, 2026