

**K941931 DIAGNOSTIC X-RAY PATIENT EXPOSURE INDICATOR**May 26, 1994  
36 days to decisionK941931 · Product code: **KPR** · Radiology  
Source: <https://www.510kdatabase.net/k941931/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k941931/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 19, 2026