

**K984111 FV-RF**Dec 11, 1998  
24 days to decisionK984111 · Product code: **KPR** · Radiology  
Source: <https://www.510kdatabase.net/k984111/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Stationary (KPR)
Date received	Nov 17, 1998
Decision date	Dec 11, 1998
Days to decision	24 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Philips Medical Systems (Cleveland), Inc.</b>
Location	Mchenry, IL, US
Contact	MICHAEL J HAYES
510(k) history	190 submissions · 190 cleared · 1977-2017

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k984111/> 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 16, 2026