

**K982196 DIGITAL RADIOGRAPHIC IMAGING SYSTEM**Oct 6, 1998  
106 days to decisionK982196 · Product code: **KPR** · Radiology  
Source: <https://www.510kdatabase.net/k982196/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Stationary (KPR)
Date received	Jun 22, 1998
Decision date	Oct 6, 1998
Days to decision	106 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>GE Medical Systems</b>
Location	Milwaukee, WI, US
Contact	LARRY A KROGER
510(k) history	169 submissions · 166 cleared · 1989-2022

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k982196/> 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 27, 2026