

**K980220 IISYS PACS SYSTEM**Feb 13, 1998  
22 days to decisionK980220 · Product code: **LMD** · Radiology  
Source: <https://www.510kdatabase.net/k980220/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Digital Image Communications, Radiological (LMD)
Date received	Jan 22, 1998
Decision date	Feb 13, 1998
Days to decision	22 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Sterling Diagnostic Imaging, Inc.</b>
Location	Newark, DE, US
Contact	TIMOTHY W CAPEHART
510(k) history	7 submissions · 7 cleared · 1996-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k980220/> 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