

**K031535 PHILIPS EASYDIAGNOST ELEVA**Jun 17, 2003  
32 days to decisionK031535 · Product code: **KPR** · Radiology  
Source: <https://www.510kdatabase.net/k031535/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, X-ray, Stationary (KPR)
Date received	May 16, 2003
Decision date	Jun 17, 2003
Days to decision	32 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Philips Medical Systems North America Co.</b>
Location	Shelton, CT, US
Contact	LYNN T HARMER
510(k) history	24 submissions · 24 cleared · 2001-2010

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k031535/> 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 17, 2026