

**K902793 RUBELLA IGG**Jul 31, 1990  
35 days to decisionK902793 · Product code: **LFX** · Microbiology  
Source: <https://www.510kdatabase.net/k902793/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Enzyme Linked Immunoabsorbent Assay, Rubella (LFX)
Date received	Jun 26, 1990
Decision date	Jul 31, 1990
Days to decision	35 days
Third-party review	No

**APPLICANT**

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Company	<b>Pharmacia Diagnostics, Inc.</b>
Location	Columbia, MD, US
Contact	KAREN DARCY
510(k) history	7 submissions · 7 cleared · 1990-1992

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k902793/>; 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 June 29, 2026