

**K901635 RUBELLA IGG ELISA TEST**Jun 28, 1990  
80 days to decisionK901635 · Product code: LFX · Microbiology  
Source: <https://www.510kdatabase.net/k901635/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Enzyme Linked Immunoabsorbent Assay, Rubella (LFX)
Date received	Apr 9, 1990
Decision date	Jun 28, 1990
Days to decision	80 days
Third-party review	No

**APPLICANT**

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Company	<b>Gull Laboratories, Inc.</b>
Location	UT, US
Contact	RACHFORD, PHD
510(k) history	57 submissions · 57 cleared · 1977-1999

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k901635/>; 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