

K073390 DIASORIN LIASON RUBELLA IGG ASSAYNov 21, 2008
354 days to decisionK073390 · Product code: LFX · Microbiology
Source: <https://www.510kdatabase.net/k073390/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Enzyme Linked Immunoabsorbent Assay, Rubella (LFX)
Date received	Dec 3, 2007
Decision date	Nov 21, 2008
Days to decision	354 days
Third-party review	No
Summary / Statement	Summary
Other names	DIASORIN LIAISON RUBELLA IGG TRI-CONTROLS

APPLICANT

Company	DiaSorin, Inc.
Location	Ellicott City, MD, US
Contact	Mari Meyer
510(k) history	71 submissions · 70 cleared · 1998-2026

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k073390/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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