

**K922801 RESP. PANEL 1 INDIRECT IMMUNOFLUORESCENCE
ASSAY**Jan 6, 1993
210 days to decisionK922801 · Product code: **GNY** · Microbiology
Source: <https://www.510kdatabase.net/k922801/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Antisera, Fluorescent, Adenovirus 1-33 (GNY)
Date received	Jun 10, 1992
Decision date	Jan 6, 1993
Days to decision	210 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Light Diagnostics
Location	Temecula, CA, US
Contact	DALE DEMBROW
510(k) history	19 submissions · 19 cleared · 1993-1999

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k922801/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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