

**K911059 TURBIDOMETRIC STANDARD**Jun 10, 1991  
91 days to decisionK911059 · Product code: **LIE** · Microbiology  
Source: <https://www.510kdatabase.net/k911059/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Reagent/device, Inoculum Calibration (LIE)
Date received	Mar 11, 1991
Decision date	Jun 10, 1991
Days to decision	91 days
Third-party review	No

**APPLICANT**

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Company	<b>Scientific Device Laboratory, Inc.</b>
Location	Mchenry, IL, US
Contact	STEWART LIPTON
510(k) history	28 submissions · 28 cleared · 1984-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k911059/>; 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 14, 2026