

**K890984 LINISTAIN SLS**Apr 4, 1989  
36 days to decisionK890984 · Product code: **KIO** · Pathology  
Source: <https://www.510kdatabase.net/k890984/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Slide Stainer, Immersion Type (KIO)
Date received	Feb 27, 1989
Decision date	Apr 4, 1989
Days to decision	36 days
Third-party review	No

**APPLICANT**

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Company	<b>Shandon, Inc.</b>
Location	Mchenry, IL, US
Contact	CROOKHAM, BA
510(k) history	5 submissions · 5 cleared · 1978-1989

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k890984/>; 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