

**K893723 LORI-MODEL L, CANAL IN-THE-EAR INSTRUMENT**Sep 8, 1989  
114 days to decisionK893723 · Product code: **ESD** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k893723/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	May 17, 1989
Decision date	Sep 8, 1989
Days to decision	114 days
Third-party review	No

**APPLICANT**

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Company	<b>Lori Medical Laboratories, Inc.</b>
Location	Long Lake, MN, US
Contact	RICHARD R MAAS
510(k) history	8 submissions · 8 cleared · 1989-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k893723/>; 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 July 5, 2026