

**K903479 AUDIOTECH MODEL DB**Oct 9, 1990  
68 days to decisionK903479 · Product code: **ESD** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k903479/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	Aug 2, 1990
Decision date	Oct 9, 1990
Days to decision	68 days
Third-party review	No

**APPLICANT**

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Company	<b>Muskogee Regional Hearing Aid Lab</b>
Location	Muskogee, OK, US
Contact	DON LANGSTON
510(k) history	4 submissions · 4 cleared · 1990-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k903479/>; 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