

**K950963 AUDIOTECH MODEL SPI**May 25, 1995  
84 days to decisionK950963 · Product code: **ESD** · Ear, Nose, Throat  
Source: <https://www.510kdatabase.net/k950963/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	Mar 2, 1995
Decision date	May 25, 1995
Days to decision	84 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k950963/> 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