FDA 510(k) Premarket Notification

K950963 AUDIOTECH MODEL SPI

CLEARED

May 25, 1995
84 days to decision

K950963 · 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	**Muskogee Regional Hearing Aid Lab**
Location	Muskogee, OK, US
Contact	DON LANGSTON
510(k) history	4 submissions · 4 cleared · 1990-1995