FDA 510(k) Premarket Notification

K903479 AUDIOTECH MODEL DB

CLEARED

Oct 9, 1990
68 days to decision

K903479 · Product code: **ESD** · Ear, Nose, Throat
Source: https://www.510kdatabase.net/k903479/

SUBMISSION DETAILS

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

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