FDA 510(k) Premarket Notification

K913245 MODEL 80 HEARING AID DISPENSING SYSTEM

CLEARED

Oct 18, 1991
88 days to decision

K913245 · Product code: **ESD** · Ear, Nose, Throat
Source: https://www.510kdatabase.net/k913245/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	Jul 22, 1991
Decision date	Oct 18, 1991
Days to decision	88 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**3M Company**
Location	White City, OR, US
Contact	RALPH FRAVEL
Website	http://www.3m.com/
510(k) history	331 submissions · 322 cleared · 1976-2025

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k913245/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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