FDA 510(k) Premarket Notification

K935095 8260 SERIES 3M PROGRAMMABLE HEARING INSTRUMENTS

CLEARED

Dec 15, 1993
51 days to decision

K935095 · Product code: **ESD** · Ear, Nose, Throat
Source: https://www.510kdatabase.net/k935095/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	Oct 25, 1993
Decision date	Dec 15, 1993
Days to decision	51 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Statement

APPLICANT

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