FDA 510(k) Premarket Notification

K790093 MODULE PERSONAL EAR (SUPER)

CLEARED

Jan 26, 1979
10 days to decision

K790093 · Product code: **ESD** · Ear, Nose, Throat
Source: https://www.510kdatabase.net/k790093/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	Jan 16, 1979
Decision date	Jan 26, 1979
Days to decision	10 days
Third-party review	No

APPLICANT

Company	**Qualitone**
Location	Mchenry, IL, US
510(k) history	39 submissions · 39 cleared · 1976-1996