FDA 510(k) Premarket Notification

K833803 OTICON, E26P

CLEARED

Dec 8, 1983
37 days to decision

K833803 · Product code: **ESD** · Ear, Nose, Throat
Source: https://www.510kdatabase.net/k833803/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	Nov 1, 1983
Decision date	Dec 8, 1983
Days to decision	37 days
Third-party review	No

APPLICANT

Company	**Oticon Corp.**
Location	Mchenry, IL, US
510(k) history	57 submissions · 57 cleared · 1978-1996