

K833803 OTICON, E26PDec 8, 1983
37 days to decisionK833803 · Product code: **ESD** · Ear, Nose, ThroatSource: <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

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Device record: <https://www.510kdatabase.net/k833803/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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