

K821325 OTICONJun 7, 1982
34 days to decisionK821325 · Product code: **ESD** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k821325/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	May 4, 1982
Decision date	Jun 7, 1982
Days to decision	34 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/k821325/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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