

**K902559 CUSTOM IN-THE-EAR HEARING AID OTICON MODEL:
P**Jul 9, 1990
28 days to decisionK902559 · Product code: **ESD** · Ear, Nose, Throat
Source: <https://www.510kdatabase.net/k902559/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	Jun 11, 1990
Decision date	Jul 9, 1990
Days to decision	28 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k902559/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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