

K905555 A-ZILICONEJul 6, 1992
573 days to decisionK905555 · Product code: **ESD** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k905555/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	Dec 11, 1990
Decision date	Jul 6, 1992
Days to decision	573 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/k905555/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 5, 2026