

K973886 OTICON DIGISOUND COMPACTNov 4, 1997
21 days to decisionK973886 · Product code: **ESD** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k973886/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	Oct 14, 1997
Decision date	Nov 4, 1997
Days to decision	21 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Oticon, Inc.
Location	Somerset, NJ, US
Contact	PREBEN BRUNVED
510(k) history	9 submissions · 9 cleared · 1993-1997

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k973886/> 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