

K933640 WIDEX MODEL LOGO 12 (L12)Aug 23, 1993
27 days to decisionK933640 · Product code: **ESD** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k933640/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	Jul 27, 1993
Decision date	Aug 23, 1993
Days to decision	27 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Widex Hearing Aid Co., Inc.
Location	Mchenry, IL, US
Contact	RON MELTSNER
510(k) history	52 submissions · 52 cleared · 1976-2008

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k933640/> 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 June 25, 2026