

**K901642 WIDEX HEARING AID MODEL THE DISCOVERY**May 23, 1990  
44 days to decisionK901642 · Product code: **ESD** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k901642/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	Apr 9, 1990
Decision date	May 23, 1990
Days to decision	44 days
Third-party review	No

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k901642/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 25, 2026