

**K832129 HEARING AID G3H & G3T**Jul 26, 1983  
25 days to decisionK832129 · Product code: **ESD** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k832129/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Air-conduction, Prescription (ESD)
Date received	Jul 1, 1983
Decision date	Jul 26, 1983
Days to decision	25 days
Third-party review	No

**APPLICANT**

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Company	<b>Widex Hearing Aid Co., Inc.</b>
Location	Mchenry, IL, US
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/k832129/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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