

**K931322 KS 5 - CLINICAL AUDIOMETER**Aug 19, 1994  
521 days to decisionK931322 · Product code: **EWO** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k931322/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Audiometer (EWO)
Date received	Mar 16, 1993
Decision date	Aug 19, 1994
Days to decision	521 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Maico Hearing Instruments, Inc.</b>
Location	Mchenry, IL, US
Contact	LOIS DONNAY
510(k) history	58 submissions · 57 cleared · 1977-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k931322/> 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 July 5, 2026