

**K960567 HF MEGA EAR**Oct 16, 1996  
250 days to decisionK960567 · Product code: **EPF** · Ear, Nose, Throat  
Source: <https://www.510kdatabase.net/k960567/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Hearing Aid, Group And Auditory Trainer (EPF)
Date received	Feb 9, 1996
Decision date	Oct 16, 1996
Days to decision	250 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Maxi-Aids, Inc.</b>
Location	Farmingdale, NY, US
Contact	HARRIS BOSHAK
510(k) history	2 submissions · 2 cleared · 1996-2004

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k960567/> 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 28, 2026