

**K071973 EPLEY OMNIAX**Jun 20, 2008  
339 days to decisionK071973 · Product code: **LXV** · Ear, Nose, Throat  
Source: <https://www.510kdatabase.net/k071973/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Apparatus, Vestibular Analysis (LXV)
Date received	Jul 17, 2007
Decision date	Jun 20, 2008
Days to decision	339 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Vesticon</b>
Location	Portland, OR, US
Contact	CATHRYN EPLEY
510(k) history	1 submissions · 1 cleared · 2008-2008

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k071973/> 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