

**K972252 VISTA UNIVERSAL HANDPIECE**Jul 29, 1997  
43 days to decisionK972252 · Product code: **HQC** · Ophthalmic  
Source: <https://www.510kdatabase.net/k972252/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Phacofragmentation (HQC)
Date received	Jun 16, 1997
Decision date	Jul 29, 1997
Days to decision	43 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Odyssey Technologies, Inc.</b>
Location	Temple City, CA, US
Contact	GLENN A DUNKI-JACOBS
510(k) history	2 submissions · 2 cleared · 1997-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k972252/> 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 29, 2026