

**K884924 VISI-WIPE**Jan 31, 1989  
64 days to decisionK884924 · Product code: **HOF** · Ophthalmic  
Source: <https://www.510kdatabase.net/k884924/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Burr, Corneal, Manual (HOF)
Date received	Nov 28, 1988
Decision date	Jan 31, 1989
Days to decision	64 days
Third-party review	No

**APPLICANT**

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Company	<b>Visitec Co.</b>
Location	Walker, MI, US
Contact	LIAQUAT ALLARAKHIA
510(k) history	49 submissions · 49 cleared · 1979-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k884924/>; 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 May 23, 2026