

**K902857 VISI-DRAPE OPHTHALMIC DRAPES**Jul 31, 1990  
32 days to decisionK902857 · Product code: **HMY** · Ophthalmic  
Source: <https://www.510kdatabase.net/k902857/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Keratome, Battery-powered (HMY)
Date received	Jun 29, 1990
Decision date	Jul 31, 1990
Days to decision	32 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/k902857/>; 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