

**K881022 VCTS MARKETING PROGRAM SYSTEM**Apr 1, 1988  
22 days to decisionK881022 · Product code: **HOX** · Ophthalmic  
Source: <https://www.510kdatabase.net/k881022/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Chart, Visual Acuity (HOX)
Date received	Mar 10, 1988
Decision date	Apr 1, 1988
Days to decision	22 days
Third-party review	No

**APPLICANT**

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Company	<b>Vistech Consultants, Inc.</b>
Location	Walker, MI, US
Contact	DENNIS J MCCREIGHT
510(k) history	7 submissions · 7 cleared · 1984-1988

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k881022/>, 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 30, 2026