

**K803256 EYETECH-VISION SCREENING SYSTEM**Mar 4, 1981  
71 days to decisionK803256 · Product code: **HOX** · Ophthalmic  
Source: <https://www.510kdatabase.net/k803256/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Chart, Visual Acuity (HOX)
Date received	Dec 23, 1980
Decision date	Mar 4, 1981
Days to decision	71 days
Third-party review	No

**APPLICANT**

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Company	<b>Eyedex</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1981-1981

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k803256/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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