

**K781452 BAYLOW VISUAL ACUITY TESTER**Oct 24, 1978  
62 days to decisionK781452 · Product code: **HOX** · Ophthalmic  
Source: <https://www.510kdatabase.net/k781452/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Chart, Visual Acuity (HOX)
Date received	Aug 23, 1978
Decision date	Oct 24, 1978
Days to decision	62 days
Third-party review	No

**APPLICANT**

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Company	<b>Codman &amp; Shurtleff, Inc.</b>
Location	Mchenry, IL, US
510(k) history	152 submissions · 151 cleared · 1976-2020

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

Device record: <https://www.510kdatabase.net/k781452/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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