

**K861066 AR-1000 AUTOMATIC REFRACTOR**Apr 4, 1986  
15 days to decisionK861066 · Product code: **HKO** · Ophthalmic  
Source: <https://www.510kdatabase.net/k861066/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Refractometer, Ophthalmic (HKO)
Date received	Mar 20, 1986
Decision date	Apr 4, 1986
Days to decision	15 days
Third-party review	No

**APPLICANT**

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Company	<b>Nidek, Inc.</b>
Location	Mchenry, IL, US
Contact	JOHN BRATKOWSKY
510(k) history	77 submissions · 77 cleared · 1983-2005

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k861066/>; 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 19, 2026