

**K841861 CANON AUTOREF R-10**Oct 5, 1984  
154 days to decisionK841861 · Product code: **HKO** · Ophthalmic  
Source: <https://www.510kdatabase.net/k841861/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Refractometer, Ophthalmic (HKO)
Date received	May 4, 1984
Decision date	Oct 5, 1984
Days to decision	154 days
Third-party review	No

**APPLICANT**

---

Company	<b>Canon USA, Inc.</b>
Location	Mchenry, IL, US
510(k) history	48 submissions · 48 cleared · 1984-2009

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k841861/> 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 18, 2026