

**K913962 ANOMALOSCOPE QUADRANT TEST-6**Oct 23, 1991  
48 days to decisionK913962 · Product code: **HIW** · Ophthalmic  
Source: <https://www.510kdatabase.net/k913962/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Anomaloscope (HIW)
Date received	Sep 5, 1991
Decision date	Oct 23, 1991
Days to decision	48 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Lkc Technologies, Inc.</b>
Location	Mchenry, IL, US
Contact	JEROME LEIGHT
510(k) history	8 submissions · 8 cleared · 1976-2015

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k913962/> 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 4, 2026