

**K904962 EYE ARTIFICIAL CUSTOM**Aug 30, 1991  
298 days to decisionK904962 · Product code: **HQH** · Ophthalmic  
Source: <https://www.510kdatabase.net/k904962/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Eye, Artificial, Non-custom (HQH)
Date received	Nov 5, 1990
Decision date	Aug 30, 1991
Days to decision	298 days
Third-party review	No

**APPLICANT**

---

Company	<b>Midwest Eye Laboratories, Inc.</b>
Location	Minneapolis, MN, US
Contact	MICHAEL BARRETT
510(k) history	2 submissions · 2 cleared · 1991-1991

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k904962/>; 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 30, 2026