

K874252 KREDA DISCJan 27, 1988
100 days to decisionK874252 · Product code: **HOQ** · Ophthalmic
Source: <https://www.510kdatabase.net/k874252/>**SUBMISSION DETAILS**

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|-----------------------|------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Grid, Amsler (HOQ) |
| Date received | Oct 19, 1987 |
| Decision date | Jan 27, 1988 |
| Days to decision | 100 days |
| Third-party review | No |

APPLICANT

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|----------------|---|
| Company | Richmond Products, Inc. |
| Location | Boca Raton, FL, US |
| Contact | LLOYD POWELL |
| 510(k) history | 15 submissions · 15 cleared · 1987-1988 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k874252/>; Data sourced from FDA 510(k) public records (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. - Generated June 30, 2026