

K862692 KARICKHOFF DIAGNOSTIC/LASER LENSAug 21, 1986
37 days to decisionK862692 · Product code: **HKS** · Ophthalmic
Source: <https://www.510kdatabase.net/k862692/>**SUBMISSION DETAILS**

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|-----------------------|------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Prism, Gonioscopic (HKS) |
| Date received | Jul 15, 1986 |
| Decision date | Aug 21, 1986 |
| Days to decision | 37 days |
| Third-party review | No |

APPLICANT

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|----------------|---|
| Company | Ocular Instruments, Inc. |
| Location | Bellevue, WA, US |
| Contact | TAMSIN J ERICKSON |
| 510(k) history | 50 submissions · 50 cleared · 1984-2002 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k862692/>; 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 29, 2026