

K842835 BRILLETTE MODELNov 21, 1984
124 days to decisionK842835 · Product code: **HOI** · Ophthalmic
Source: <https://www.510kdatabase.net/k842835/>**SUBMISSION DETAILS**

| | |
|-----------------------|------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Spectacle, Magnifying (HOI) |
| Date received | Jul 20, 1984 |
| Decision date | Nov 21, 1984 |
| Days to decision | 124 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---------------------------------------|
| Company | Cillus Company, Inc. |
| Location | Novato, CA, US |
| Contact | JORGEN L KREJSBOL |
| 510(k) history | 1 submissions · 1 cleared · 1984-1984 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k842835/>; 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 July 1, 2026