

K900279 OCULINUM INJECTION AMPLIFIERFeb 9, 1990
32 days to decisionK900279 · Product code: **HLW** · Ophthalmic
Source: <https://www.510kdatabase.net/k900279/>**SUBMISSION DETAILS**

| | |
|-----------------------|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Preamplifier, Battery-powered, Ophthalmic (HLW) |
| Date received | Jan 8, 1990 |
| Decision date | Feb 9, 1990 |
| Days to decision | 32 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---------------------------------------|
| Company | Oculinum, Inc. |
| Location | Mill Valley, CA, US |
| Contact | SCOTT, M.D. |
| 510(k) history | 3 submissions · 2 cleared · 1986-1990 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k900279/>; 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