

K890992 WOODLYN OPHTHALMIC PROJECTORMay 17, 1989
79 days to decisionK890992 · Product code: **HOS** · Ophthalmic
Source: <https://www.510kdatabase.net/k890992/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Projector, Ophthalmic (HOS)
Date received	Feb 27, 1989
Decision date	May 17, 1989
Days to decision	79 days
Third-party review	No

APPLICANT

Company	Woodlyn, Inc.
Location	Mchenry, IL, US
Contact	RONALD NIELSEN
510(k) history	19 submissions · 19 cleared · 1984-1993

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k890992/>; 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