FDA 510(k) Premarket Notification

K900279 OCULINUM INJECTION AMPLIFIER

CLEARED

Feb 9, 1990
32 days to decision

K900279 · 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