FDA 510(k) Premarket Notification

K896072 D.L. SCOPE OPHTHALMOSCOPE

CLEARED

Nov 27, 1989
39 days to decision

K896072 · Product code: **HLJ** · Ophthalmic
Source: https://www.510kdatabase.net/k896072/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ophthalmoscope, Battery-powered (HLJ)
Date received	Oct 19, 1989
Decision date	Nov 27, 1989
Days to decision	39 days
Third-party review	No

APPLICANT

Company	**North American Medical Products, Inc.**
Location	Walker, MI, US
Contact	ARTHUR GIANAKOS
510(k) history	6 submissions · 6 cleared · 1984-2001

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k896072/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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