

K844364 SKIASCOPIIC RACKSJan 2, 1985
50 days to decisionK844364 · Product code: **HMH** · Ophthalmic
Source: <https://www.510kdatabase.net/k844364/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Rack, Skiascopic (HMH)
Date received	Nov 13, 1984
Decision date	Jan 2, 1985
Days to decision	50 days
Third-party review	No

APPLICANT

Company	Luneau Laboratories
Location	Washington, DC, US
510(k) history	5 submissions · 5 cleared · 1984-1985

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k844364/>; 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