FDA 510(k) Premarket Notification

K844364 SKIASCOPIC RACKS

CLEARED

Jan 2, 1985
50 days to decision

K844364 · 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