

K801767 DIMENSION I REFRACTION SYSTEMSep 16, 1980
50 days to decisionK801767 · Product code: **HLM** · Ophthalmic
Source: <https://www.510kdatabase.net/k801767/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Measuring, Lens, Ac-powered (HLM)
Date received	Jul 28, 1980
Decision date	Sep 16, 1980
Days to decision	50 days
Third-party review	No

APPLICANT

Company	Smr
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1980-1981

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Device record: <https://www.510kdatabase.net/k801767/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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