

K802183 SR-IV PROGRAMMED SUBJECTIVE REFRACTORApr 23, 1981
226 days to decisionK802183 · Product code: **HKO** · Ophthalmic
Source: <https://www.510kdatabase.net/k802183/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Refractometer, Ophthalmic (HKO)
Date received	Sep 9, 1980
Decision date	Apr 23, 1981
Days to decision	226 days
Third-party review	No

APPLICANT

Company	American Optical Corp.
Location	Mchenry, IL, US
510(k) history	35 submissions · 35 cleared · 1976-1995

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Device record: <https://www.510kdatabase.net/k802183/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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