

K801906 DIOPTRON III AUTOREFRACTORAug 27, 1980
16 days to decisionK801906 · Product code: **HKO** · Ophthalmic
Source: <https://www.510kdatabase.net/k801906/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Refractometer, Ophthalmic (HKO)
Date received	Aug 11, 1980
Decision date	Aug 27, 1980
Days to decision	16 days
Third-party review	No

APPLICANT

Company	Coherent Medical Division
Location	Mchenry, IL, US
510(k) history	6 submissions · 6 cleared · 1979-1990

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Device record: <https://www.510kdatabase.net/k801906/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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