

K821083 P.D. METERJun 14, 1982
56 days to decisionK821083 · Product code: **HLH** · Ophthalmic
Source: <https://www.510kdatabase.net/k821083/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pupillometer, Manual (HLH)
Date received	Apr 19, 1982
Decision date	Jun 14, 1982
Days to decision	56 days
Third-party review	No

APPLICANT

Company	Medical Equipment Designs, Inc.
Location	Mchenry, IL, US
510(k) history	13 submissions · 13 cleared · 1982-1991

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Device record: <https://www.510kdatabase.net/k821083/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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