FDA 510(k) Premarket Notification

K821083 P.D. METER

CLEARED

Jun 14, 1982
56 days to decision

K821083 · 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