FDA 510(k) Premarket Notification

K801767 DIMENSION I REFRACTION SYSTEM

CLEARED

Sep 16, 1980
50 days to decision

K801767 · 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