FDA 510(k) Premarket Notification

K861746 KEELER DUAL LIGHTSOURCE

CLEARED

Aug 21, 1986
107 days to decision

K861746 · Product code: **HPQ** · Ophthalmic
Source: https://www.510kdatabase.net/k861746/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Headlamp, Operating, Ac-powered (HPQ)
Date received	May 6, 1986
Decision date	Aug 21, 1986
Days to decision	107 days
Third-party review	No

APPLICANT

Company	**Keeler Instruments, Inc.**
Location	Mchenry, IL, US
Contact	VAN ARSDALE
510(k) history	60 submissions · 60 cleared · 1981-2019