FDA 510(k) Premarket Notification

K862692 KARICKHOFF DIAGNOSTIC/LASER LENS

CLEARED

Aug 21, 1986
37 days to decision

K862692 · Product code: **HKS** · Ophthalmic
Source: https://www.510kdatabase.net/k862692/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prism, Gonioscopic (HKS)
Date received	Jul 15, 1986
Decision date	Aug 21, 1986
Days to decision	37 days
Third-party review	No

APPLICANT

Company	**Ocular Instruments, Inc.**
Location	Bellevue, WA, US
Contact	TAMSIN J ERICKSON
510(k) history	50 submissions · 50 cleared · 1984-2002