FDA 510(k) Premarket Notification

K832939 I-GLIDE IOL IMPLANTATION AID

CLEARED

Oct 4, 1983
35 days to decision

K832939 · Product code: **KYB** · Ophthalmic
Source: https://www.510kdatabase.net/k832939/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lens, Guide, Intraocular (KYB)
Date received	Aug 30, 1983
Decision date	Oct 4, 1983
Days to decision	35 days
Third-party review	No

APPLICANT

Company	**I For M, Inc.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1983-1983