FDA 510(k) Premarket Notification

K821381 IOL FORCEPS

CLEARED

Jun 14, 1982
35 days to decision

K821381 · Product code: **KYB** · Ophthalmic
Source: https://www.510kdatabase.net/k821381/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lens, Guide, Intraocular (KYB)
Date received	May 10, 1982
Decision date	Jun 14, 1982
Days to decision	35 days
Third-party review	No

APPLICANT

Company	**Cilco, Inc.**
Location	Mchenry, IL, US
510(k) history	12 submissions · 12 cleared · 1982-1986