FDA 510(k) Premarket Notification

K883489 KRATZ NOVA SERT(TM)

CLEARED

Oct 25, 1988
70 days to decision

K883489 · Product code: **KYB** · Ophthalmic
Source: https://www.510kdatabase.net/k883489/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lens, Guide, Intraocular (KYB)
Date received	Aug 16, 1988
Decision date	Oct 25, 1988
Days to decision	70 days
Third-party review	No

APPLICANT

Company	**Coopervision-Cilco**
Location	Bellevue, WA, US
Contact	RAY NEGLAY
510(k) history	2 submissions · 2 cleared · 1988-1988