FDA 510(k) Premarket Notification

K902489 MODIFIED MOLTENO(TM) IMPLANTS

CLEARED

Aug 14, 1990
90 days to decision

K902489 · Product code: **KYF** · Ophthalmic
Source: https://www.510kdatabase.net/k902489/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Eye Valve (KYF)
Date received	May 16, 1990
Decision date	Aug 14, 1990
Days to decision	90 days
Third-party review	No

APPLICANT

Company	**Iop, Inc.**
Location	Malibu, CA, US
Contact	JASON MALECKA
510(k) history	9 submissions · 8 cleared · 1989-2009