FDA 510(k) Premarket Notification

K890300 INTERCASSETTE, INTERKIT, UNIVERSAL PAK & CASSETTE

CLEARED

Apr 17, 1989
88 days to decision

K890300 · Product code: **HQE** · Ophthalmic
Source: https://www.510kdatabase.net/k890300/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Vitreous Aspiration And Cutting, Ac-powered (HQE)
Date received	Jan 19, 1989
Decision date	Apr 17, 1989
Days to decision	88 days
Third-party review	No

APPLICANT

Company	**CooperVision, Inc.**
Location	Southampton, GB
Contact	DAVE KRAPF
Website	https://www.coopervision.com
510(k) history	97 submissions · 94 cleared · 1978-2024

CooperVision, Inc. is a contact lens manufacturer based in Southampton, GB. The company specializes in ophthalmic devices for vision correction. CooperVision has received FDA 510(k) clearances from total submissions since its first clearance in 1978. Ophthalmic devices represent 88% of the company's regulatory submissions. The company remains active, with its latest FDA 510(k) clearance in 2024. Recent cleared devices include daily disposable contact lenses in spheric, toric, and multifocal designs. Notable product families include MyDay, Clariti 1 day, Biofinity, and Ava...