

**K854375 COOPERVISION ACAP POWER DRIVE ULTRASONIC ANTERIOR**Dec 27, 1985  
58 days to decisionK854375 · Product code: **HQC** · Ophthalmic  
Source: <https://www.510kdatabase.net/k854375/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Phacofragmentation (HQC)
Date received	Oct 30, 1985
Decision date	Dec 27, 1985
Days to decision	58 days
Third-party review	No

**APPLICANT**

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Company	<b>CooperVision, Inc.</b>
Location	Southampton, GB
Contact	DAVID W KRAPF
Website	<a href="https://www.coopervision.com">https://www.coopervision.com</a>
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...

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