

**K890635 CAVITRON/KELMAN PHACOEMULSIFIER  
ASPIRATOR, UNIV.**May 9, 1989  
90 days to decisionK890635 · Product code: **HQC** · Ophthalmic  
Source: <https://www.510kdatabase.net/k890635/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Phacofragmentation (HQC)
Date received	Feb 8, 1989
Decision date	May 9, 1989
Days to decision	90 days
Third-party review	No

**APPLICANT**

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Company	<b>CooperVision, Inc.</b>
Location	Southampton, GB
Contact	DAVE 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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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k890635/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 28, 2026