

**K831836 EPSILON ULTRASONIC TIP**Jun 30, 1983  
23 days to decisionK831836 · Product code: **HQC** · Ophthalmic  
Source: <https://www.510kdatabase.net/k831836/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Phacofragmentation (HQC)
Date received	Jun 7, 1983
Decision date	Jun 30, 1983
Days to decision	23 days
Third-party review	No

**APPLICANT**

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Company	<b>CooperVision, Inc.</b>
Location	Southampton, GB
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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