

**K832142 DICON GLAUCOMA DIAG. CENTER 2000 SERIES**Aug 1, 1983  
27 days to decisionK832142 · Product code: **HPT** · Ophthalmic  
Source: <https://www.510kdatabase.net/k832142/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Perimeter, Automatic, Ac-powered (HPT)
Date received	Jul 5, 1983
Decision date	Aug 1, 1983
Days to decision	27 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 spherical, toric, and multifocal designs. Notable product families include MyDay, Clariti 1 day, Biofinity, and Ava...

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