

**K843039 SITE ANTERIOR CAPSULOTOMER**Nov 21, 1984  
111 days to decisionK843039 · Product code: **HMX** · Ophthalmic  
Source: <https://www.510kdatabase.net/k843039/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Ophthalmic (HMX)
Date received	Aug 2, 1984
Decision date	Nov 21, 1984
Days to decision	111 days
Third-party review	No

**APPLICANT**

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Company	<b>Chiron Vision Corp.</b>
Location	Irvine, CA, US
Contact	TICKNOR
510(k) history	34 submissions · 34 cleared · 1980-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k843039/>; 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 29, 2026