

**K883489 KRATZ NOVA SERT(TM)**Oct 25, 1988  
70 days to decisionK883489 · Product code: **KYB** · Ophthalmic  
Source: <https://www.510kdatabase.net/k883489/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lens, Guide, Intraocular (KYB)
Date received	Aug 16, 1988
Decision date	Oct 25, 1988
Days to decision	70 days
Third-party review	No

**APPLICANT**

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Company	<b>Coopervision-Cilco</b>
Location	Bellevue, WA, US
Contact	RAY NEGLAY
510(k) history	2 submissions · 2 cleared · 1988-1988

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k883489/>; 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 May 21, 2026