

**K862070 INTER-SHARP**Jun 17, 1986  
18 days to decisionK862070 · Product code: **HNW** · Ophthalmic  
Source: <https://www.510kdatabase.net/k862070/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dilator, Lachrymal (HNW)
Date received	May 30, 1986
Decision date	Jun 17, 1986
Days to decision	18 days
Third-party review	No

**APPLICANT**

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Company	<b>Intermedics Intraocular, Inc.</b>
Location	Walker, MI, US
Contact	ENID ROKAW
510(k) history	4 submissions · 4 cleared · 1983-1986

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k862070/>, 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 30, 2026