

**K902489 MODIFIED MOLTENO(TM) IMPLANTS**Aug 14, 1990  
90 days to decisionK902489 · Product code: **KYF** · Ophthalmic  
Source: <https://www.510kdatabase.net/k902489/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Eye Valve (KYF)
Date received	May 16, 1990
Decision date	Aug 14, 1990
Days to decision	90 days
Third-party review	No

**APPLICANT**

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Company	<b>Iop, Inc.</b>
Location	Malibu, CA, US
Contact	JASON MALECKA
510(k) history	9 submissions · 8 cleared · 1989-2009

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k902489/>; 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 23, 2026