

**K832939 I-GLIDE IOL IMPLANTATION AID**Oct 4, 1983  
35 days to decisionK832939 · Product code: **KYB** · Ophthalmic  
Source: <https://www.510kdatabase.net/k832939/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lens, Guide, Intraocular (KYB)
Date received	Aug 30, 1983
Decision date	Oct 4, 1983
Days to decision	35 days
Third-party review	No

**APPLICANT**

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Company	<b>I For M, Inc.</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1983-1983

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k832939/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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