

**K023737 INJECTOR**Jul 14, 2003  
249 days to decisionK023737 · Product code: **MSS** · Ophthalmic  
Source: <https://www.510kdatabase.net/k023737/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Folders And Injectors, Intraocular Lens (iol) (MSS)
Date received	Nov 7, 2002
Decision date	Jul 14, 2003
Days to decision	249 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Anton Meyer &amp; Co. , Ltd.</b>
Location	Nidau, CH
Contact	THOMAS MEYER
510(k) history	1 submissions · 1 cleared · 2003-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k023737/> 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