

**K931839 ROTA SYSTEM XL**Nov 22, 1993  
223 days to decisionK931839 · Product code: **HMF** · Ophthalmic  
Source: <https://www.510kdatabase.net/k931839/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stand, Instrument, Ac-powered, Ophthalmic (HMF)
Date received	Apr 13, 1993
Decision date	Nov 22, 1993
Days to decision	223 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Rota Systems Medfurniture, Inc.</b>
Location	Derby, KS, US
Contact	SAM R MYERS
510(k) history	2 submissions · 2 cleared · 1993-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k931839/> 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