

**K934880 MTI PHOTOSCREENER**Apr 6, 1994  
176 days to decisionK934880 · Product code: **MMF** · Ophthalmic  
Source: <https://www.510kdatabase.net/k934880/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Photorefractor (MMF)
Date received	Oct 12, 1993
Decision date	Apr 6, 1994
Days to decision	176 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Medical Technology and Innovations, Inc.</b>
Location	Grand Prairie, TX, US
Contact	JEREMY P FEAKINS
510(k) history	3 submissions · 3 cleared · 1986-2002

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k934880/> 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