

**K000224 FLUOPERM 60-OK, PARAGON HDS-OK**Apr 17, 2000  
84 days to decisionK000224 · Product code: **MUW** · Ophthalmic  
Source: <https://www.510kdatabase.net/k000224/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - U
Submission type	Traditional
Device classification	Lens, Contact (orthokeratology) (MUW)
Date received	Jan 24, 2000
Decision date	Apr 17, 2000
Days to decision	84 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Paragon Vision Sciences</b>
Location	Phoenix, AZ, US
Contact	WILLIAM E MEYERS
510(k) history	15 submissions · 13 cleared · 1994-2018

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k000224/> 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 May 31, 2026