

**K022560 M2 SINGLE USE MICROKERTOME**Sep 27, 2002  
56 days to decisionK022560 · Product code: **HMY** · Ophthalmic  
Source: <https://www.510kdatabase.net/k022560/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Keratome, Battery-powered (HMY)
Date received	Aug 2, 2002
Decision date	Sep 27, 2002
Days to decision	56 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Moria SA</b>
Location	Antony, FR
Contact	MELANIE RENAUD-SAMIRI
510(k) history	5 submissions · 5 cleared · 2002-2017

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k022560/> 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