

**K020482 KERATOME BLADE 200200 & 600600**Dec 16, 2002  
306 days to decisionK020482 · Product code: **HNO** · Ophthalmic  
Source: <https://www.510kdatabase.net/k020482/>**SUBMISSION DETAILS**

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|                       |                                    |
|-----------------------|------------------------------------|
| Decision              | Substantially Equivalent (Cleared) |
| Submission type       | Traditional                        |
| Device classification | Keratome, Ac-powered (HNO)         |
| Date received         | Feb 13, 2002                       |
| Decision date         | Dec 16, 2002                       |
| Days to decision      | 306 days                           |
| Third-party review    | No                                 |
| Summary / Statement   | Summary                            |

**APPLICANT**

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|----------------|---------------------------------------|
| Company        | <b>Micro Specialties, Inc.</b>        |
| Location       | Milford, CT, US                       |
| Contact        | CHARLES VASSALLO                      |
| 510(k) history | 4 submissions · 4 cleared · 1998-2004 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k020482/> 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