

**K960395 PLANCON MIRCOLAMELLAR KERATOME**Apr 26, 1996  
88 days to decisionK960395 · Product code: **HNO** · Ophthalmic  
Source: <https://www.510kdatabase.net/k960395/>**SUBMISSION DETAILS**

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|                       |                                    |
|-----------------------|------------------------------------|
| Decision              | Substantially Equivalent (Cleared) |
| Submission type       | Traditional                        |
| Device classification | Keratome, Ac-powered (HNO)         |
| Date received         | Jan 29, 1996                       |
| Decision date         | Apr 26, 1996                       |
| Days to decision      | 88 days                            |
| Third-party review    | No                                 |
| Summary / Statement   | Summary                            |

**APPLICANT**

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|----------------|---------------------------------------|
| Company        | <b>Plancon Instruments</b>            |
| Location       | Antony, FR                            |
| Contact        | ALAIN DUPRAT                          |
| 510(k) history | 3 submissions · 3 cleared · 1996-1998 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k960395/> 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