

**K924780 BURTON OPHTHALMIC EXAM UNIT - MODEL 2201**Apr 5, 1993  
194 days to decisionK924780 · Product code: **HMF** · Ophthalmic  
Source: <https://www.510kdatabase.net/k924780/>**SUBMISSION DETAILS**

---

|                       |   |
|-----------------------|---|
| Decision              | Substantially Equivalent (Cleared)              |
| Submission type       | Traditional                                     |
| Device classification | Stand, Instrument, Ac-powered, Ophthalmic (HMF) |
| Date received         | Sep 23, 1992                                    |
| Decision date         | Apr 5, 1993                                     |
| Days to decision      | 194 days  |
| Third-party review    | No  |
| Summary / Statement   | Statement                                       |

**APPLICANT**

---

|                |   |
|----------------|---|
| Company        | <b>R.H. Burton Co.</b>                  |
| Location       | Grove City, OH, US                      |
| Contact        | KEVIN M LOYCHIK                         |
| 510(k) history | 19 submissions · 19 cleared · 1992-1993 |

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k924780/> 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 22, 2026