

**K920571 TRIAL LENS SET**May 7, 1992  
90 days to decisionK920571 · Product code: **HPC** · Ophthalmic  
Source: <https://www.510kdatabase.net/k920571/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Lens, Trial, Ophthalmic (HPC)
Date received	Feb 7, 1992
Decision date	May 7, 1992
Days to decision	90 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k920571/> 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