

**K894720 UNIVERSAL EYESHIELD**Sep 14, 1989  
51 days to decisionK894720 · Product code: **HOY** · Ophthalmic  
Source: <https://www.510kdatabase.net/k894720/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Shield, Eye, Ophthalmic (including Sunlamp Protective Eyewear And Post-mydratic Eyewear) (HOY)
Date received	Jul 25, 1989
Decision date	Sep 14, 1989
Days to decision	51 days
Third-party review	No

**APPLICANT**

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Company	<b>Trident Medical Products, Inc.</b>
Location	Fort Worth, TX, US
Contact	DAN E BRUHL
510(k) history	3 submissions · 3 cleared · 1989-1989

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k894720/>, 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 30, 2026