

**K932300 FOG CONTROL, LENS FOG REDUCTION DEVICE**Sep 17, 1993  
128 days to decisionK932300 · Product code: **OCT** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k932300/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Anti Fog Solution And Accessories, Endoscopy (OCT)
Date received	May 12, 1993
Decision date	Sep 17, 1993
Days to decision	128 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Highpoint Medical Corp.</b>
Location	Arlington, TX, US
Contact	ROGER LIEBELT
510(k) history	2 submissions · 2 cleared · 1991-1993

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k932300/> 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 July 2, 2026