

**K933369 CUDA PRODUCTS CORP. LIGHTSOURCES**Jan 3, 1994  
178 days to decisionK933369 · Product code: **FCW** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k933369/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light Source, Fiberoptic, Routine (FCW)
Date received	Jul 9, 1993
Decision date	Jan 3, 1994
Days to decision	178 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Cuda Products Co.</b>
Location	Jacksonville, FL, US
Contact	CYNTHIA ARCUSA
510(k) history	17 submissions · 17 cleared · 1988-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k933369/> 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