

**K003577 STERIS DEEPSITE FIBER OPTIC SURGICAL LIGHT**Jan 29, 2001  
70 days to decisionK003577 · Product code: **FST** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k003577/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Light, Surgical, Fiberoptic (FST)
Date received	Nov 20, 2000
Decision date	Jan 29, 2001
Days to decision	70 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>STERIS Corporation</b>
Location	Mentor, OH, US
Contact	RAYMOND URSICK
510(k) history	204 submissions · 202 cleared · 1997-2026

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k003577/> 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 28, 2026