

**K083710 RETRACTOR LITE**Feb 23, 2009  
70 days to decisionK083710 · Product code: **FST** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k083710/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Surgical, Fiberoptic (FST)
Date received	Dec 15, 2008
Decision date	Feb 23, 2009
Days to decision	70 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Thomas Surgical Instruments, Inc.</b>
Location	Traverse City, MI, US
Contact	STEPHANIE A ZALUCHA
510(k) history	1 submissions · 1 cleared · 2009-2009

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k083710/> 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 23, 2026