

**K122637 LITE DECOMPRESSION SYSTEM- LIGHT CABLE
MODEL 48089999**Nov 1, 2012
64 days to decisionK122637 · Product code: FST · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k122637/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Light, Surgical, Fiberoptic (FST)
Date received	Aug 29, 2012
Decision date	Nov 1, 2012
Days to decision	64 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Stryker Spine
Location	Allendale, NJ, US
Contact	TINA MORNAK
510(k) history	74 submissions · 73 cleared · 2004-2026

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k122637/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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