FDA 510(k) Premarket Notification

K122637 LITE DECOMPRESSION SYSTEM- LIGHT CABLE MODEL 48089999

CLEARED

Nov 1, 2012
64 days to decision

K122637 · 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