

**K241171 Spine Guidance Software (Version 5.1)**Jul 25, 2024  
90 days to decisionK241171 · Product code: **OLO** · Orthopedic  
Source: <https://www.510kdatabase.net/k241171/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Orthopedic Stereotaxic Instrument (OLO)
Date received	Apr 26, 2024
Decision date	Jul 25, 2024
Days to decision	90 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary
Other names	Consolidated Operating Room Equipment (CORE) 2 Console; CORE Q Footswitch; PI Drive 2 Motors; Elite Q Attachments and Cutting Accessories

**APPLICANT**

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Company	<b>Stryker Leibinger GmbH &amp; Co KG</b>
Location	Freiburg Im Breisgau, DE
Contact	John Chesney
510(k) history	21 submissions · 21 cleared · 2009-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k241171/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://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](https://accessdata.fda.gov). - Generated May 13, 2026