

K172034 Stryker Spine Navigation System with SpineMap 3D software application, OrthoLock, nGenius Spine Clamp, Navigated Drill Guide Set, Navigated Xia 3 Awl Tap, Navigated Xia 3 Serrato TapOct 18, 2017
105 days to decisionK172034 · Product code: **OLO** · Orthopedic
Source: <https://www.510kdatabase.net/k172034/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Orthopedic Stereotaxic Instrument (OLO)
Date received	Jul 5, 2017
Decision date	Oct 18, 2017
Days to decision	105 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Stryker Corporation
Location	Malwah, NJ, US
Contact	Andrea Wallen-Gerding
Website	http://www.stryker.com/
510(k) history	81 submissions · 81 cleared · 2010-2023

Stryker Corporation is an American multinational medical technology company headquartered in Portage, Michigan. The company develops and markets surgical equipment, neurotechnology, orthopedic implants, and patient safety systems used globally across medical specialties. Stryker has received FDA 510(k) clearances from total submissions between 2010 and 2023. The company's cleared devices span orthopedic surgery, neurosurgery, general and plastic surgery, and ear, nose, and throat specialties. This regulatory record reflects the company's broad portfolio across surgical an...

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