

**K120434 POWER ADAPTOR INSTRUMENT ACCESSORY**Mar 8, 2012  
24 days to decisionK120434 · Product code: **NKB** · Orthopedic  
Source: <https://www.510kdatabase.net/k120434/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	Feb 13, 2012
Decision date	Mar 8, 2012
Days to decision	24 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Stryker Corp.</b>
Location	Mchenry, IL, US
Contact	TIFFANI ROGERS
510(k) history	124 submissions · 121 cleared · 1976-2023

Stryker Corp. is an American multinational medical technology company headquartered in Portage, Michigan. The company develops and markets surgical equipment, implants, and patient safety technologies used globally across multiple medical specialties. Stryker has received FDA 510(k) clearances from total submissions since its first clearance in 1976. The company maintains active regulatory engagement, with its latest clearance in 2023. Its product portfolio spans orthopedic devices, neurosurgical implants, surgical instruments, and endoscopy systems, reflecting a broad pr...

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