

**K203319 SYMPHONY™ OCT System**Feb 5, 2021  
85 days to decisionK203319 · Product code: **NKG** · Orthopedic  
Source: <https://www.510kdatabase.net/k203319/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Posterior Cervical Screw System (NKG)
Date received	Nov 12, 2020
Decision date	Feb 5, 2021
Days to decision	85 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Medos International SARL</b>
Location	Raynham, MA, US
Contact	Eric Zhu
510(k) history	96 submissions · 96 cleared · 2010-2026

**REGULATORY CONSULTANT**

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Consulting firm	<b>Depuy Synthes Spine</b>
Contact	Eric Zhu

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k203319/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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