

K233366 SYMPHONY OCT SystemNov 9, 2023
38 days to decisionK233366 · Product code: **NKG** · Orthopedic
Source: <https://www.510kdatabase.net/k233366/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Posterior Cervical Screw System (NKG)
Date received	Oct 2, 2023
Decision date	Nov 9, 2023
Days to decision	38 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Medos International SARL
Location	Raynham, MA, US
Contact	Sheree Geller
510(k) history	96 submissions · 96 cleared · 2010-2026

REGULATORY CONSULTANT

Consulting firm	Depuy Synthes Spine
Contact	Sheree Geller

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k233366/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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