

# K254157 CROSSNAV Navigation Enabled Instruments for Disc Preparation and Interbody

May 26, 2026  
155 days to decision

K254157 · Product code: **OLO** · Orthopedic  
Source: <https://www.510kdatabase.net/k254157/>

## SUBMISSION DETAILS

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Orthopedic Stereotaxic Instrument (OLO)
Date received	Dec 22, 2025
Decision date	May 26, 2026
Days to decision	155 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	Marina Minnock
510(k) history	97 submissions · 97 cleared · 2010-2026

## REGULATORY CONSULTANT

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Consulting firm	<b>Synthes GmbH</b>
Contact	Marina Minnock

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

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Device record: <https://www.510kdatabase.net/k254157/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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