

K252873 Q Interbody InstrumentsFeb 6, 2026
149 days to decisionK252873 · Product code: **OLO** · Orthopedic
Source: <https://www.510kdatabase.net/k252873/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Orthopedic Stereotaxic Instrument (OLO)
Date received	Sep 10, 2025
Decision date	Feb 6, 2026
Days to decision	149 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	K2m, Inc.
Location	Leesburg, VA, US
Contact	Renee Maciag
510(k) history	100 submissions · 97 cleared · 2007-2026

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k252873/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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