

K131425 NAVIGATED CAPSTONE TRIALS, CLYDESDALE TRIALS, CAPSTONE & CLYDESDALE INSERTERAug 14, 2013
89 days to decisionK131425 · Product code: **OLO** · Orthopedic
Source: <https://www.510kdatabase.net/k131425/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Orthopedic Stereotaxic Instrument (OLO)
Date received	May 17, 2013
Decision date	Aug 14, 2013
Days to decision	89 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Medtronic Navigation, Inc.
Location	Lafayette, CO, US
Contact	REGINA HOLMES
Website	https://www.medtronic.com
510(k) history	35 submissions · 35 cleared · 2005-2026

Medtronic Navigation, Inc. is based in Lafayette, US. The company specializes in surgical navigation and imaging systems for minimally invasive procedures. The company has received FDA 510(k) clearances from total submissions since 2005. Neurology devices represent a dominant category, including cranial and laser ablation systems. The latest clearance in 2026 reflects continued regulatory activity and product innovation. Recent cleared devices span multiple surgical specialties. StealthStation and Stealth AXiS™ platforms serve neurology, orthopedic, and ear, nose, and thr...

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Device record: <https://www.510kdatabase.net/k131425/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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