

K221928 ROSA® Knee SystemOct 4, 2022
95 days to decisionK221928 · Product code: **OLO** · Orthopedic
Source: <https://www.510kdatabase.net/k221928/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Orthopedic Stereotaxic Instrument (OLO)
Date received	Jul 1, 2022
Decision date	Oct 4, 2022
Days to decision	95 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Orthosoft D/B/A Zimmer Cas
Location	Montreal, CA
Contact	Kavina Veeren
510(k) history	7 submissions · 7 cleared · 2021-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k221928/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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