

K220522 7D Surgical System - Percutaneous Application (7D Flash Frame)

May 20, 2022
86 days to decision

K220522 · Product code: **OLO** · Orthopedic
Source: <https://www.510kdatabase.net/k220522/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Orthopedic Stereotaxic Instrument (OLO)
Date received	Feb 23, 2022
Decision date	May 20, 2022
Days to decision	86 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	7D Surgical
Location	Toronto, CA
Contact	Daniel Ziskind
510(k) history	1 submissions · 1 cleared · 2022-2022

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k220522/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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