

K234150 Lucent® XPMar 27, 2024
89 days to decisionK234150 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k234150/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Dec 29, 2023
Decision date	Mar 27, 2024
Days to decision	89 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Spinal Elements
Location	Carlsbad, CA, US
Contact	Julie Lamothe
510(k) history	1 submissions · 1 cleared · 2024-2024

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k234150/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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