

K251741 PathLoc Lumbar Interbody Fusion Cage SystemJul 8, 2025
32 days to decisionK251741 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k251741/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Jun 6, 2025
Decision date	Jul 8, 2025
Days to decision	32 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	L & K Biomed Co., Ltd.
Location	Yongin-Si, KR
Contact	Kihayng Kim
Website	https://www.lkbiomed.com
510(k) history	54 submissions · 54 cleared · 2010-2026

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k251741/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 13, 2026