FDA 510(k) Premarket Notification

K251940 PathLoc Lumbar Plate System

CLEARED

Mar 3, 2026
252 days to decision

K251940 · Product code: **KWQ** · Orthopedic
Source: https://www.510kdatabase.net/k251940/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Intervertebral Body (KWQ)
Date received	Jun 24, 2025
Decision date	Mar 3, 2026
Days to decision	252 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	Kihyang Kim
Website	https://www.lkbiomed.com
510(k) history	54 submissions · 54 cleared · 2010-2026