

**K251940 PathLoc Lumbar Plate System**Mar 3, 2026  
252 days to decisionK251940 · 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	<b>L &amp; K Biomed Co., Ltd.</b>
Location	Yongin-Si, KR
Contact	Kihyang Kim
Website	<a href="https://www.lkbiomed.com">https://www.lkbiomed.com</a>
510(k) history	54 submissions · 54 cleared · 2010-2026

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k251940/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://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](https://accessdata.fda.gov). - Generated May 14, 2026