

K200066 Posterior Fusion Plate, HA Posterior Fusion PlateDec 10, 2020
332 days to decisionK200066 · Product code: **PEK** · Orthopedic
Source: <https://www.510kdatabase.net/k200066/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spinous Process Plate (PEK)
Date received	Jan 13, 2020
Decision date	Dec 10, 2020
Days to decision	332 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Spinal Simplicity, LLC
Location	Philedelphia, PA, US
Contact	Adam Rogers
510(k) history	14 submissions · 14 cleared · 2015-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k200066/> 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 26, 2026