

**K001080 SPINE SYSTEM EVOLUTION**Jun 29, 2000  
86 days to decisionK001080 · Product code: **KWP** · Orthopedic  
Source: <https://www.510kdatabase.net/k001080/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Interlaminar (KWP)
Date received	Apr 4, 2000
Decision date	Jun 29, 2000
Days to decision	86 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Aesculap, Inc.</b>
Location	Burlingame, CA, US
Contact	LIA S JONES
510(k) history	207 submissions · 201 cleared · 1991-2025

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k001080/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 21, 2026