

**K191424 Interphalangeal Joint Fusion Device Range**Aug 28, 2019  
91 days to decisionK191424 · Product code: **HWC** · Orthopedic  
Source: <https://www.510kdatabase.net/k191424/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Screw, Fixation, Bone (HWC)
Date received	May 29, 2019
Decision date	Aug 28, 2019
Days to decision	91 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Neoste</b>
Location	Round Rock, TX, US
Contact	Anne-Cecile Grandremy
510(k) history	6 submissions · 6 cleared · 2013-2022

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k191424/> 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 27, 2026