

**K200312 Tryptik Ti**Apr 6, 2020  
60 days to decisionK200312 · Product code: **ODP** · Orthopedic  
Source: <https://www.510kdatabase.net/k200312/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Cervical (ODP)
Date received	Feb 6, 2020
Decision date	Apr 6, 2020
Days to decision	60 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Spineart</b>
Location	Geneva, CH
Contact	Frank Pennesi
510(k) history	44 submissions · 44 cleared · 2008-2023

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k200312/> 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 14, 2026