

**K233223 OGMend® Implant Enhancement System**Oct 26, 2023  
28 days to decisionK233223 · Product code: **QVI** · Orthopedic  
Source: <https://www.510kdatabase.net/k233223/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Screw Sleeve Bone Fixation Device, Spine (QVI)
Date received	Sep 28, 2023
Decision date	Oct 26, 2023
Days to decision	28 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Woven Orthopedic Technologies</b>
Location	Manchester, CT, US
Contact	Brandon Bendes
510(k) history	2 submissions · 2 cleared · 2023-2023

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k233223/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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