

K173153 Vertera Spine™ Coalesce™ Thoracolumbar Interbody Fusion SystemDec 6, 2017
68 days to decisionK173153 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k173153/>**SUBMISSION DETAILS**

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|-----------------------|--|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Intervertebral Fusion Device With Bone Graft, Lumbar (MAX) |
| Date received | Sep 29, 2017 |
| Decision date | Dec 6, 2017 |
| Days to decision | 68 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---|
| Company | Nu Vasive, Incorporated |
| Location | San Diego, CA, US |
| Contact | Michelle Cheung |
| 510(k) history | 112 submissions · 112 cleared · 2012-2023 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k173153/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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