

K172275 Vitality® Spinal Fixation System, Vitality®+ Power Instrument System

Sep 27, 2017
61 days to decision

K172275 · Product code: **NKB** · Orthopedic
Source: <https://www.510kdatabase.net/k172275/>

SUBMISSION DETAILS

| | |
|-----------------------|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Thoracolumbosacral Pedicle Screw System (NKB) |
| Date received | Jul 28, 2017 |
| Decision date | Sep 27, 2017 |
| Days to decision | 61 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---|
| Company | Zimmer Biomet Spine, Inc. |
| Location | Broomfield, CO, US |
| Contact | Kelly Stratton |
| 510(k) history | 15 submissions · 15 cleared · 2017-2021 |

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k172275/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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