

K172469 TiLock Modular Spinal SystemNov 30, 2017
107 days to decisionK172469 · Product code: **NKB** · Orthopedic
Source: <https://www.510kdatabase.net/k172469/>**SUBMISSION DETAILS**

| | |
|-----------------------|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Thoracolumbosacral Pedicle Screw System (NKB) |
| Date received | Aug 15, 2017 |
| Decision date | Nov 30, 2017 |
| Days to decision | 107 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---|
| Company | Genesys Spine |
| Location | Austin, TX, US |
| Contact | Brian J. Bergeron |
| 510(k) history | 31 submissions · 31 cleared · 2010-2025 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k172469/> Data sourced from FDA 510(k) public records (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. - Generated May 16, 2026