

K170512 Altus Spine Titanium Interbody Fusion SystemMar 10, 2017
17 days to decisionK170512 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k170512/>**SUBMISSION DETAILS**

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|-----------------------|--|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Special |
| Device classification | Intervertebral Fusion Device With Bone Graft, Lumbar (MAX) |
| Date received | Feb 21, 2017 |
| Decision date | Mar 10, 2017 |
| Days to decision | 17 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---|
| Company | Altus Partners, LLC |
| Location | Newtown Square, PA, US |
| Contact | Claudia Hill |
| 510(k) history | 17 submissions · 17 cleared · 2015-2025 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k170512/> 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 26, 2026