

K120401 ALAMO TOct 5, 2012
240 days to decisionK120401 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k120401/>**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 | Feb 8, 2012 |
| Decision date | Oct 5, 2012 |
| Days to decision | 240 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---|
| Company | Alliance Partners, LLC |
| Location | Avon Lake, OH, US |
| Contact | JENNIFER PALINCHIK |
| 510(k) history | 10 submissions · 10 cleared · 2011-2020 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k120401/> 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 June 18, 2026