

K130573 TYBER MEDICAL INTERBODY SYSTEMSep 30, 2013
210 days to decisionK130573 · Product code: **ODP** · Orthopedic
Source: <https://www.510kdatabase.net/k130573/>**SUBMISSION DETAILS**

| | |
|-----------------------|--|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Intervertebral Fusion Device With Bone Graft, Cervical (ODP) |
| Date received | Mar 4, 2013 |
| Decision date | Sep 30, 2013 |
| Days to decision | 210 days |
| Third-party review | No |
| Summary / Statement | Summary |
| Other names | ACIF, ALIF, PLIF, TLIF, DLIF |

APPLICANT

| | |
|----------------|---|
| Company | Tyber Medical, LLC |
| Location | Morristown, NJ, US |
| Contact | JEFF TYBER |
| Website | https://www.tybermed.com |
| 510(k) history | 28 submissions · 28 cleared · 2013-2026 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k130573/> 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 13, 2026