

K173864 Peekplus® Interbody Fusion DeviceJul 2, 2018
194 days to decisionK173864 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k173864/>**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 | Dec 20, 2017 |
| Decision date | Jul 2, 2018 |
| Days to decision | 194 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---------------------------------------|
| Company | Vallum Corporation |
| Location | Nashua, NH, US |
| Contact | Stephen Blinn |
| 510(k) history | 1 submissions · 1 cleared · 2018-2018 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k173864/> 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