

K181261 Curiteva Cervical Interbody Fusion SystemJul 9, 2018
59 days to decisionK181261 · Product code: **ODP** · Orthopedic
Source: <https://www.510kdatabase.net/k181261/>**SUBMISSION DETAILS**

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|-----------------------|--|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Intervertebral Fusion Device With Bone Graft, Cervical (ODP) |
| Date received | May 11, 2018 |
| Decision date | Jul 9, 2018 |
| Days to decision | 59 days |
| Third-party review | No |
| Combination product | No |
| PCCP authorized | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---------------------------------------|
| Company | Curiteva, LLC |
| Location | Tanner, AL, US |
| Contact | Eric Linder |
| 510(k) history | 4 submissions · 4 cleared · 2018-2018 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k181261/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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