

K073348 LUCENT MAGNUMDec 26, 2007
27 days to decisionK073348 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k073348/>**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 | Nov 29, 2007 |
| Decision date | Dec 26, 2007 |
| Days to decision | 27 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---|
| Company | Spinal Elements, Inc. |
| Location | Carlsbad, CA, US |
| Contact | KERRI DIMARTINO |
| Website | https://www.spinalelements.com |
| 510(k) history | 48 submissions · 48 cleared · 2007-2026 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k073348/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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