FDA 510(k) Premarket Notification

# K131077 LIFE SPINE PLATEAU SPACER SYSTEM

**CLEARED**

Jul 17, 2013
91 days to decision

K131077 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k131077/

## 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 | Apr 17, 2013 |
| Decision date | Jul 17, 2013 |
| Days to decision | 91 days |
| Third-party review | No |
| Summary / Statement | Summary |

## APPLICANT

| | |
|---|---|
| Company | **Life Spine** |
| Location | Hoffman Estates, IL, US |
| Contact | RANDY LEWIS |
| 510(k) history | 36 submissions · 34 cleared · 2006-2018 |