FDA 510(k) Premarket Notification

# K143126 Renovis S141 Lumbar Interbody Cage System

**CLEARED**

Feb 6, 2015
98 days to decision

K143126 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k143126/

## 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 | Oct 31, 2014 |
| Decision date | Feb 6, 2015 |
| Days to decision | 98 days |
| Third-party review | No |
| Summary / Statement | Summary |

## APPLICANT

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|---|---|
| Company | **Renovis Surgical Technologies** |
| Location | Redlands, CA, US |
| Contact | Josh Brown |
| 510(k) history | 7 submissions · 7 cleared · 2015-2019 |