FDA 510(k) Premarket Notification

# K172130 Resorbable Mesh Device

**CLEARED**

Nov 16, 2017
125 days to decision

K172130 · Product code: **MQV** · Orthopedic
Source: https://www.510kdatabase.net/k172130/

## SUBMISSION DETAILS

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|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Filler, Bone Void, Calcium Compound (MQV) |
| Date received | Jul 14, 2017 |
| Decision date | Nov 16, 2017 |
| Days to decision | 125 days |
| Third-party review | No |
| Summary / Statement | Summary |

## APPLICANT

| | |
|---|---|
| Company | **SeaSpine Orthopedics Corporation** |
| Location | Carlsbad, CA, US |
| Contact | Jenny Fam |
| 510(k) history | 66 submissions · 66 cleared · 2016-2025 |