

K173260 SeaSpine Spacer System (NanoMetalene)- Hollywood, Hollywood VI, Ventura, PacificaDec 4, 2017
55 days to decisionK173260 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k173260/>**SUBMISSION DETAILS**

| | |
|-----------------------|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Intervertebral Fusion Device With Bone Graft, Lumbar (MAX) |
| Date received | Oct 10, 2017 |
| Decision date | Dec 4, 2017 |
| Days to decision | 55 days |
| Third-party review | No |
| Summary / Statement | Summary |
| Other names | SeaSpine Spacer System -Hollywood, Hollywood VI, Ventura, Pacifica; SeaSpine Vu e-POD System |

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k173260/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 15, 2026