FDA 510(k) Premarket Notification

# K102020 VEGA SPAN SPINOUS PROCESS PLATE SYSTEM

**CLEARED**

Sep 1, 2010
44 days to decision

K102020 · Product code: **KWP** · Orthopedic
Source: https://www.510kdatabase.net/k102020/

## SUBMISSION DETAILS

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|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Appliance, Fixation, Spinal Interlaminal (KWP) |
| Date received | Jul 19, 2010 |
| Decision date | Sep 1, 2010 |
| Days to decision | 44 days |
| Third-party review | No |
| Summary / Statement | Summary |

## APPLICANT

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|---|---|
| Company | **Spinefrontier, Inc.** |
| Location | Beverly, MA, US |
| Contact | JOHN SULLIVAN |
| 510(k) history | 24 submissions · 24 cleared · 2007-2020 |