

K173702 Juliet Ti LLJan 3, 2018
30 days to decisionK173702 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k173702/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Dec 4, 2017
Decision date	Jan 3, 2018
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Spineart
Location	Geneva, CH
Contact	Franck Pennesi
510(k) history	44 submissions · 44 cleared · 2008-2023

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