

K172392 Lateral Spinal Truss System (LSTS) Interbody Fusion DeviceNov 7, 2017
91 days to decisionK172392 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k172392/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Aug 8, 2017
Decision date	Nov 7, 2017
Days to decision	91 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	4Web, Inc.
Location	Frisco, TX, US
Contact	Jessee Hunt
510(k) history	13 submissions · 13 cleared · 2014-2026

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k172392/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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