FDA 510(k) Premarket Notification

CLEARED

K172392 Lateral Spinal Truss System (LSTS) Interbody Fusion Device

Nov 7, 2017
91 days to decision

K172392 · 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