

K143126 Renovis S141 Lumbar Interbody Cage SystemFeb 6, 2015
98 days to decisionK143126 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k143126/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Oct 31, 2014
Decision date	Feb 6, 2015
Days to decision	98 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Renovis Surgical Technologies
Location	Redlands, CA, US
Contact	Josh Brown
510(k) history	7 submissions · 7 cleared · 2015-2019

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k143126/> 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 June 28, 2026