

K170888 Renovis S141 Lumbar Interbody Fusion SystemAug 8, 2017
134 days to decisionK170888 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k170888/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Mar 27, 2017
Decision date	Aug 8, 2017
Days to decision	134 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k170888/> 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