

K183447 EIT Cellular Titanium Lumbar Cage - T/PLIFFeb 4, 2019
54 days to decisionK183447 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k183447/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Dec 12, 2018
Decision date	Feb 4, 2019
Days to decision	54 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Eit Emerging Implant Technologies GmbH
Location	Wurmlingen, DE
Contact	Barbara Wirth
510(k) history	5 submissions · 5 cleared · 2017-2020

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k183447/> 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