

K162327 COUGAR® LS Lateral Cage System and COUGAR® System

Dec 12, 2016
115 days to decision

K162327 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k162327/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Aug 19, 2016
Decision date	Dec 12, 2016
Days to decision	115 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Medos International SARL
Location	Raynham, MA, US
Contact	Eric Zhu
510(k) history	96 submissions · 96 cleared · 2010-2026

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k162327/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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