

K071983 AESCULAP PEEK INTERVERTEBRAL BODY FUSION SYSTEM: A SPACE, CSPACE, PROSPACE, T-SPACEMay 1, 2008
287 days to decisionK071983 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k071983/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Jul 19, 2007
Decision date	May 1, 2008
Days to decision	287 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Aesculap Implant Systems, Inc.
Location	Center Valley, PA, US
Contact	MATTHEW M HULL
510(k) history	22 submissions · 22 cleared · 2007-2016

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k071983/> 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 May 17, 2026