

K143479 Bluefin Interbody SystemSep 25, 2015
291 days to decisionK143479 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k143479/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Dec 8, 2014
Decision date	Sep 25, 2015
Days to decision	291 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Tides Medical
Location	Lafayette, LA, US
Contact	J. Doug Payne
510(k) history	1 submissions · 1 cleared · 2015-2015

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k143479/> 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