

**K122021 STERISPINE LC CAGE**Sep 20, 2012  
72 days to decisionK122021 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k122021/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Jul 10, 2012
Decision date	Sep 20, 2012
Days to decision	72 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Safe Orthopaedics</b>
Location	Eragny Sur Oise, FR
Contact	P. DUMOUCHEL
510(k) history	13 submissions · 13 cleared · 2011-2020

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k122021/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 18, 2026