

**K250186 GoLIF! Lumbar Interbody Fusion System**Mar 21, 2025  
58 days to decisionK250186 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k250186/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Jan 22, 2025
Decision date	Mar 21, 2025
Days to decision	58 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Getset Surgical, SA</b>
Location	Epalinges, CH
Contact	Jowita Sokalska
510(k) history	2 submissions · 2 cleared · 2021-2025

**REGULATORY CONSULTANT**

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Consulting firm	<b>MRC Global</b>
Contact	Christine Scifert

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k250186/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://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](https://accessdata.fda.gov). - Generated May 14, 2026