

**K231809 Momentum® Posterior Spinal Fixation System with G21 V-STEADY Bone Cement**Oct 20, 2023  
122 days to decisionK231809 · Product code: **PML** · Orthopedic  
Source: <https://www.510kdatabase.net/k231809/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bone Cement, Posterior Screw Augmentation (PML)
Date received	Jun 20, 2023
Decision date	Oct 20, 2023
Days to decision	122 days
Third-party review	No
Summary / Statement	Summary
Other names	Momentum® MIS Posterior Spinal Fixation System with G21 V-STEADY Bone Cement

**APPLICANT**

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Company	<b>Ulrich Medical USA</b>
Location	Chesterfield, MD, US
Contact	Eric Lucas
510(k) history	8 submissions · 8 cleared · 2015-2023

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k231809/> 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 25, 2026