

**K180090 Cure™ Lumbar Plate System**Apr 23, 2018  
101 days to decisionK180090 · Product code: **KWQ** · Orthopedic  
Source: <https://www.510kdatabase.net/k180090/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Appliance, Fixation, Spinal Intervertebral Body (KWQ)
Date received	Jan 12, 2018
Decision date	Apr 23, 2018
Days to decision	101 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Meditech Spine, LLC</b>
Location	Atlanta, GA, US
Contact	Bruce Dunaway
510(k) history	10 submissions · 10 cleared · 2015-2021

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k180090/> 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 May 27, 2026