

**K223303 Spinery™ RF Ablation System**Aug 30, 2023  
307 days to decisionK223303 · Product code: **GEI** · Orthopedic  
Source: <https://www.510kdatabase.net/k223303/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Oct 27, 2022
Decision date	Aug 30, 2023
Days to decision	307 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Axon Spine Medical System</b>
Location	Pompei, IT
Contact	Salvatore Accardo
510(k) history	1 submissions · 1 cleared · 2023-2023

**REGULATORY CONSULTANT**

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Consulting firm	<b>Quality Solutions and Support, LLC</b>
Contact	Tawney Schwarz

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k223303/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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