

**K091228 SPINEWAND SURGICAL DEVICE**Sep 28, 2009  
154 days to decisionK091228 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k091228/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Apr 27, 2009
Decision date	Sep 28, 2009
Days to decision	154 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Arthrocare Corp.</b>
Location	Mountain View, CA, US
Contact	VALERIE DEFIESTA-NG
Website	<a href="http://www.arthrocare.com/">http://www.arthrocare.com/</a>
510(k) history	112 submissions · 112 cleared · 1995-2016

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k091228/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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