

**K072089 ARTHROCARE SPINEWAND**Aug 17, 2007  
18 days to decisionK072089 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k072089/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jul 30, 2007
Decision date	Aug 17, 2007
Days to decision	18 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/k072089/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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