

**K100353 SPINEWAND SURGICAL DEVICE**May 13, 2010  
90 days to decisionK100353 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k100353/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Feb 12, 2010
Decision date	May 13, 2010
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

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

---

**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k100353/> Data sourced from FDA 510(k) public records (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 14, 2026