

**K132099 ARTHROCARE COBLATOR IQ DLR SPINEWAND,  
ARTHROCARE COBLATOR IQ DLG SPINEWAND**Aug 22, 2013  
45 days to decisionK132099 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k132099/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jul 8, 2013
Decision date	Aug 22, 2013
Days to decision	45 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>ArthroCare Corporation</b>
Location	Irvine, CA, US
Contact	SHIRLEY HYINK
Website	<a href="http://www.arthrocare.com/">http://www.arthrocare.com/</a>
510(k) history	36 submissions · 36 cleared · 2011-2026

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k132099/> 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 June 8, 2026