

**K014290 ENTEC PLASMA WANDS**

Mar 28, 2002  
90 days to decision

K014290 · Product code: **GEI** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k014290/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Dec 28, 2001
Decision date	Mar 28, 2002
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Arthrocare Corp.</b>
Location	Mountain View, CA, US
Contact	BRUCE PROTHRO
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/k014290/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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