

K021364 MODIFICATION TO ENTEC PLASMA WANDSMay 30, 2002
30 days to decisionK021364 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k021364/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Apr 30, 2002
Decision date	May 30, 2002
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k021364/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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