

**K022475 MODIFICATION TO ARTHROCARE CONTROLLER
(SYSTEM 2000 AND 8000), ARTHROCARE PATIENT CABLE,
FOOT CONTROL, POWER CORD, WANDS**Oct 3, 2002
66 days to decisionK022475 · Product code: **GEI** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k022475/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jul 29, 2002
Decision date	Oct 3, 2002
Days to decision	66 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/k022475/> 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. - Generated May 14, 2026