

K020622 ARTHROCARE CONTROLLER (SYSTEM 2000 AND 8000)

Mar 28, 2002
30 days to decision

K020622 · Product code: **GEI** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k020622/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Feb 26, 2002
Decision date	Mar 28, 2002
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary
Other names	ARTHROCARE CABLE; FOOT CONTROL; POWER CORD; AND WANDS

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.net

Device record: <https://www.510kdatabase.net/k020622/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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