FDA 510(k) Premarket Notification

K030551 ARTHROCARE CONTROLLER (SYSTEM 2000 AND 8000)

CLEARED

Mar 7, 2003
14 days to decision

K030551 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k030551/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Feb 21, 2003
Decision date	Mar 7, 2003
Days to decision	14 days
Third-party review	No
Summary / Statement	Summary
Other names	ARTHROCARE PATIENT CABLE; FOOT CONTROL; POWER CORD; 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/k030551/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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