

**K001302 ARTHROCARE SYSTEM 2000 CONTROLLER,
ARTHROCARE SYSTEM 2000 CABLE, ARTHROCARE SYSTEM
2000 FOOTSWITCH, PLASMA SCALPEL GS, C**

May 30, 2000
36 days to decision

K001302 · Product code: **GEI** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k001302/>

SUBMISSION DETAILS

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|-----------------------|--|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Electrosurgical, Cutting & Coagulation & Accessories (GEI) |
| Date received | Apr 24, 2000 |
| Decision date | May 30, 2000 |
| Days to decision | 36 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---|
| Company | Arthrocare Corp. |
| Location | Mountain View, CA, US |
| Contact | BRUCE PROTHRO |
| 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/k001302/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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