

**K102496 POWERHEART AED G3 SEMI-AUTOMATIC,
POWERHEART AED G3 AUTOMATIC, POWERHEART AED G3
PRO, POWERHEART AED G3 PLUS**Jun 9, 2011
282 days to decisionK102496 · Product code: **MKJ** · Cardiovascular
Source: <https://www.510kdatabase.net/k102496/>**SUBMISSION DETAILS**

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|-----------------------|--|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Special |
| Device classification | Automated External Defibrillators (non-wearable) (MKJ) |
| Date received | Aug 31, 2010 |
| Decision date | Jun 9, 2011 |
| Days to decision | 282 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---|
| Company | Cardiac Science Corporation |
| Location | Bothell, WA, US |
| Contact | KATHLEEN ROBERTS |
| 510(k) history | 10 submissions · 10 cleared · 2006-2015 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k102496/> 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 June 4, 2026