

K192715 CRF Radiofrequency Ablation SystemJan 7, 2020
102 days to decisionK192715 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k192715/>**SUBMISSION DETAILS**

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|-----------------------|--|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Electrosurgical, Cutting & Coagulation & Accessories (GEI) |
| Date received | Sep 27, 2019 |
| Decision date | Jan 7, 2020 |
| Days to decision | 102 days |
| Third-party review | No |
| Combination product | No |
| PCCP authorized | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---------------------------------------|
| Company | Cambridge Interventional, LLC |
| Location | Burlington, MA, US |
| Contact | Michael A. Arnold |
| 510(k) history | 1 submissions · 1 cleared · 2020-2020 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k192715/> 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 29, 2026