

**K042388 RADIANT MEDICAL ENDOCATHETER TEMPERATURE  
PROBE**Oct 28, 2004  
56 days to decisionK042388 · Product code: **DQY** · Cardiovascular  
Source: <https://www.510kdatabase.net/k042388/>**SUBMISSION DETAILS**

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|                       |                                    |
|-----------------------|------------------------------------|
| Decision              | Substantially Equivalent (Cleared) |
| Submission type       | Traditional                        |
| Device classification | Catheter, Percutaneous (DQY)       |
| Date received         | Sep 2, 2004                        |
| Decision date         | Oct 28, 2004                       |
| Days to decision      | 56 days                            |
| Third-party review    | No                                 |
| Summary / Statement   | Summary                            |

**APPLICANT**

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|----------------|---------------------------------------|
| Company        | <b>Radiant Medical</b>                |
| Location       | Redwood City, CA, US                  |
| Contact        | ANDREW CLEELAND                       |
| 510(k) history | 6 submissions · 5 cleared · 2002-2007 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k042388/> 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](http://accessdata.fda.gov). - Generated June 28, 2026