

**K070539 MORTARA ELI 10 ELECTROCARDIOGRAPH**Mar 15, 2007  
17 days to decisionK070539 · Product code: **DPS** · Cardiovascular  
Source: <https://www.510kdatabase.net/k070539/>**SUBMISSION DETAILS**

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|                       |                                    |
|-----------------------|------------------------------------|
| Decision              | Substantially Equivalent (Cleared) |
| Submission type       | Special                            |
| Device classification | Electrocardiograph (DPS)           |
| Date received         | Feb 26, 2007                       |
| Decision date         | Mar 15, 2007                       |
| Days to decision      | 17 days                            |
| Third-party review    | No                                 |
| Summary / Statement   | Summary                            |

**APPLICANT**

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|----------------|---|
| Company        | <b>Mortara Instrument, Inc.</b>         |
| Location       | Walker, MI, US                          |
| Contact        | LAURA M SPIEGELHOFF                     |
| 510(k) history | 51 submissions · 51 cleared · 1983-2019 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k070539/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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 19, 2026