

**K041548 SENTEC DIGITAL MONITOR MODEL #SDM**Aug 9, 2004  
61 days to decisionK041548 · Product code: **LKD** · Anesthesiology  
Source: <https://www.510kdatabase.net/k041548/>**SUBMISSION DETAILS**

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|                       |  |
|-----------------------|--|
| Decision              | Substantially Equivalent (Cleared)       |
| Submission type       | Traditional                              |
| Device classification | Monitor, Carbon-dioxide, Cutaneous (LKD) |
| Date received         | Jun 9, 2004                              |
| Decision date         | Aug 9, 2004                              |
| Days to decision      | 61 days                                  |
| Third-party review    | No                                       |
| Summary / Statement   | Summary                                  |

**APPLICANT**

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|----------------|---------------------------------------|
| Company        | <b>Sentec, Inc.</b>                   |
| Location       | Egale, WI, US                         |
| Contact        | STEPHEN GORSKI                        |
| 510(k) history | 1 submissions · 1 cleared · 2004-2004 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k041548/> 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 May 25, 2026