

**K992227 CRIT-LINE MONITOR III (CLM III)**Jul 30, 1999  
28 days to decisionK992227 · Product code: **MQS** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k992227/>**SUBMISSION DETAILS**

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|                       |   |
|-----------------------|---|
| Decision              | Substantially Equivalent (Cleared)                          |
| Submission type       | Special   |
| Device classification | System, Hemodialysis, Access Recirculation Monitoring (MQS) |
| Date received         | Jul 2, 1999   |
| Decision date         | Jul 30, 1999  |
| Days to decision      | 28 days   |
| Third-party review    | No  |
| Summary / Statement   | Summary   |

**APPLICANT**

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|----------------|---------------------------------------|
| Company        | <b>In-Line Diagnostics Corp.</b>      |
| Location       | Riverdale, UT, US                     |
| Contact        | MATTHEW L HAYNIE                      |
| 510(k) history | 8 submissions · 8 cleared · 1995-2000 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k992227/> 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 28, 2026