

**K944120 FI02 SENSING MODULE**May 10, 1995  
260 days to decisionK944120 · Product code: **CCL** · Anesthesiology  
Source: <https://www.510kdatabase.net/k944120/>**SUBMISSION DETAILS**

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|                       |  |
|-----------------------|--|
| Decision              | Substantially Equivalent (Cleared)         |
| Submission type       | Traditional                                |
| Device classification | Analyzer, Gas, Oxygen, Gaseous-phase (CCL) |
| Date received         | Aug 23, 1994                               |
| Decision date         | May 10, 1995                               |
| Days to decision      | 260 days                                   |
| Third-party review    | No   |
| Summary / Statement   | Statement                                  |

**APPLICANT**

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|----------------|---------------------------------------|
| Company        | <b>Core-M, L.P.</b>                   |
| Location       | Allston, MA, US                       |
| Contact        | NATAN E PARSONS                       |
| 510(k) history | 1 submissions · 1 cleared · 1995-1995 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k944120/> 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 26, 2026