

**K991112 SAGE CONTRAST MANAGEMENT SYSTEM**Sep 2, 1999  
154 days to decisionK991112 · Product code: **FPA** · General HospitalSource: <https://www.510kdatabase.net/k991112/>**SUBMISSION DETAILS**

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|-----------------------|--|
| Decision              | Substantially Equivalent (Cleared)       |
| Submission type       | Traditional                              |
| Device classification | Set, Administration, Intravascular (FPA) |
| Date received         | Apr 1, 1999                              |
| Decision date         | Sep 2, 1999                              |
| Days to decision      | 154 days                                 |
| Third-party review    | No                                       |
| Summary / Statement   | Statement                                |

**APPLICANT**

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|----------------|---------------------------------------|
| Company        | <b>Sage Laboratories, Inc.</b>        |
| Location       | Shreveport, LA, US                    |
| Contact        | DAVID CHEN                            |
| 510(k) history | 1 submissions · 1 cleared · 1999-1999 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k991112/> 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 July 4, 2026