

**K020944 VORTEX VALVED HOLDING CHAMBER**May 9, 2002  
45 days to decisionK020944 · Product code: **CAF** · Anesthesiology  
Source: <https://www.510kdatabase.net/k020944/>**SUBMISSION DETAILS**

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|                       |                                            |
|-----------------------|--------------------------------------------|
| Decision              | Substantially Equivalent (Cleared)         |
| Submission type       | Traditional                                |
| Device classification | Nebulizer (direct Patient Interface) (CAF) |
| Date received         | Mar 25, 2002                               |
| Decision date         | May 9, 2002                                |
| Days to decision      | 45 days                                    |
| Third-party review    | Yes                                        |
| Summary / Statement   | Summary                                    |

**APPLICANT**

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|----------------|--------------------------------------------|
| Company        | <b>Pari Innovative Manufacturers, Inc.</b> |
| Location       | Midlothian, VA, US                         |
| Contact        | LAWRENCE WEINSTEIN                         |
| 510(k) history | 10 submissions · 10 cleared · 2002-2007    |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k020944/> 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 24, 2026