

**K962042 GLASS SYRINGE**Oct 8, 1996  
133 days to decisionK962042 · Product code: **FMF** · General Hospital  
Source: <https://www.510kdatabase.net/k962042/>**SUBMISSION DETAILS**

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|                       |                                    |
|-----------------------|------------------------------------|
| Decision              | Substantially Equivalent (Cleared) |
| Submission type       | Traditional                        |
| Device classification | Syringe, Piston (FMF)              |
| Date received         | May 28, 1996                       |
| Decision date         | Oct 8, 1996                        |
| Days to decision      | 133 days                           |
| Third-party review    | No                                 |
| Summary / Statement   | Statement                          |

**APPLICANT**

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|----------------|---------------------------------------|
| Company        | <b>Ispg, Inc.</b>                     |
| Location       | New Milford, CT, US                   |
| Contact        | LORETTA LUHMAN                        |
| 510(k) history | 4 submissions · 4 cleared · 1995-2001 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k962042/> 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