

**K954823 VENIPUNCTURE KIT**Jan 30, 1996  
102 days to decisionK954823 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k954823/>**SUBMISSION DETAILS**

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|                       |  |
|-----------------------|--|
| Decision              | Substantially Equivalent (Cleared)       |
| Submission type       | Traditional                              |
| Device classification | Set, Administration, Intravascular (FPA) |
| Date received         | Oct 20, 1995                             |
| Decision date         | Jan 30, 1996                             |
| Days to decision      | 102 days                                 |
| Third-party review    | No                                       |
| Summary / Statement   | Statement                                |

**APPLICANT**

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|----------------|---------------------------------------|
| Company        | <b>Winfield Industries</b>            |
| Location       | San Diego, CA, US                     |
| Contact        | DAVID OLMSTEAD                        |
| 510(k) history | 3 submissions · 3 cleared · 1994-1996 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k954823/> 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 6, 2026