

**K923882 IV-TUBE SECURE**Dec 24, 1992  
142 days to decisionK923882 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k923882/>**SUBMISSION DETAILS**

---

|                       |  |
|-----------------------|--|
| Decision              | Substantially Equivalent (Cleared)       |
| Submission type       | Traditional                              |
| Device classification | Set, Administration, Intravascular (FPA) |
| Date received         | Aug 4, 1992                              |
| Decision date         | Dec 24, 1992                             |
| Days to decision      | 142 days                                 |
| Third-party review    | No                                       |
| Summary / Statement   | Statement                                |

**APPLICANT**

---

|                |                                       |
|----------------|---------------------------------------|
| Company        | <b>Ximedix, Inc.</b>                  |
| Location       | Colorado Springs, CO, US              |
| Contact        | FRANK D SULARZ                        |
| 510(k) history | 1 submissions · 1 cleared · 1992-1992 |

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k923882/> 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 7, 2026