

**K951702 FMS HIGH FLOW ARTHRO-SHEATH**May 12, 1995  
30 days to decisionK951702 · Product code: **HNX** · Ophthalmic  
Source: <https://www.510kdatabase.net/k951702/>**SUBMISSION DETAILS**

---

|                       |                               |
|-----------------------|-------------------------------|
| Decision              | Substantially Equivalent - SN |
| Submission type       | Traditional                   |
| Device classification | Depressor, Orbital (HNX)      |
| Date received         | Apr 12, 1995                  |
| Decision date         | May 12, 1995                  |
| Days to decision      | 30 days                       |
| Third-party review    | No                            |

**APPLICANT**

---

|                |   |
|----------------|---|
| Company        | <b>Future Medical Systems, Inc.</b>     |
| Location       | New York City, NY, US                   |
| Contact        | PATRICK JANIN                           |
| 510(k) history | 12 submissions · 11 cleared · 1995-2003 |

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k951702/>; 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 14, 2026