

K910173 SELECTRA AND TUBING SETSJan 13, 1992
363 days to decisionK910173 · Product code: **FKX** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k910173/>**SUBMISSION DETAILS**

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|-----------------------|--|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | System, Peritoneal, Automatic Delivery (FKX) |
| Date received | Jan 15, 1991 |
| Decision date | Jan 13, 1992 |
| Days to decision | 363 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---------------------------------------|
| Company | Medionics International , Ltd. |
| Location | Mchenry, IL, US |
| Contact | MAHESH AGARWAL |
| 510(k) history | 6 submissions · 6 cleared · 1982-1992 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k910173/>; Data sourced from FDA 510(k) public records (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. - Generated July 2, 2026