

K893909 MODIFIED MATERIAL USED IN INFUSION TAug 25, 1989
87 days to decisionK893909 · Product code: **FPA** · General HospitalSource: <https://www.510kdatabase.net/k893909/>**SUBMISSION DETAILS**

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| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Set, Administration, Intravascular (FPA) |
| Date received | May 30, 1989 |
| Decision date | Aug 25, 1989 |
| Days to decision | 87 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---|
| Company | Quinton, Inc. |
| Location | Mchenry, IL, US |
| Contact | RON R DUCK |
| 510(k) history | 164 submissions · 160 cleared · 1976-2003 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k893909/>; 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 May 20, 2026