

**K896252 MODIFIED QUINTON SINGLE LUMEN CATHETER**Jan 29, 1990  
91 days to decisionK896252 · Product code: **MPB** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k896252/>**SUBMISSION DETAILS**

---

|                       |   |
|-----------------------|---|
| Decision              | Substantially Equivalent (Cleared)          |
| Submission type       | Traditional                                 |
| Device classification | Catheter, Hemodialysis, Non-implanted (MPB) |
| Date received         | Oct 30, 1989                                |
| Decision date         | Jan 29, 1990                                |
| Days to decision      | 91 days                                     |
| Third-party review    | No  |

**APPLICANT**

---

|                |   |
|----------------|---|
| Company        | <b>Quinton, Inc.</b>                      |
| Location       | Mchenry, IL, US                           |
| Contact        | RANDY WALLS                               |
| 510(k) history | 164 submissions · 160 cleared · 1976-2003 |

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k896252/>; 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 20, 2026