

K143135 Spring Coil GuidewireFeb 20, 2015
112 days to decisionK143135 · Product code: **DQX** · Cardiovascular
Source: <https://www.510kdatabase.net/k143135/>**SUBMISSION DETAILS**

| | |
|-----------------------|------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Wire, Guide, Catheter (DQX) |
| Date received | Oct 31, 2014 |
| Decision date | Feb 20, 2015 |
| Days to decision | 112 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---|
| Company | Neometrics, Inc. |
| Location | Mchenry, IL, US |
| Contact | Gene Champeau |
| 510(k) history | 21 submissions · 21 cleared · 1978-2015 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k143135/> 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 June 3, 2026