

**K010511 MODIFICATION TO: AGILITY STEERABLE
GUIDEWIRES**Mar 1, 2001
8 days to decisionK010511 · Product code: **DQX** · Cardiovascular
Source: <https://www.510kdatabase.net/k010511/>**SUBMISSION DETAILS**

| | |
|-----------------------|------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Special |
| Device classification | Wire, Guide, Catheter (DQX) |
| Date received | Feb 21, 2001 |
| Decision date | Mar 1, 2001 |
| Days to decision | 8 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---|
| Company | Cordis Neurovascular, Inc. |
| Location | Miami Lakes, FL, US |
| Contact | Amarilys Machado |
| 510(k) history | 37 submissions · 37 cleared · 1994-2008 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k010511/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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