

**K121776 AGILITY STEERABLE GUIDEWIRE NEUROSCOUT  
STEERABLE GUDIEWIRE**Aug 14, 2012  
57 days to decisionK121776 · Product code: **DQX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k121776/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Jun 18, 2012
Decision date	Aug 14, 2012
Days to decision	57 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Codman &amp; Shurtleff, Inc.</b>
Location	Mchenry, IL, US
Contact	JOAN BARTLE
510(k) history	152 submissions · 151 cleared · 1976-2020

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k121776/> 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 15, 2026