

**K120229 ENVOY DISTAL ACCESS GUIDING CATHETER**Feb 24, 2012  
30 days to decisionK120229 · Product code: **DQY** · CardiovascularSource: <https://www.510kdatabase.net/k120229/>**SUBMISSION DETAILS**

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|                       |                                    |
|-----------------------|------------------------------------|
| Decision              | Substantially Equivalent (Cleared) |
| Submission type       | Special                            |
| Device classification | Catheter, Percutaneous (DQY)       |
| Date received         | Jan 25, 2012                       |
| Decision date         | Feb 24, 2012                       |
| Days to decision      | 30 days                            |
| Third-party review    | No                                 |
| Summary / Statement   | Summary                            |

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k120229/> 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 June 15, 2026