

**K000715 ENVOY AND VISTA BRITE TIP**Mar 31, 2000  
29 days to decisionK000715 · Product code: **DQY** · CardiovascularSource: <https://www.510kdatabase.net/k000715/>**SUBMISSION DETAILS**

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|                       |                                    |
|-----------------------|------------------------------------|
| Decision              | Substantially Equivalent (Cleared) |
| Submission type       | Special                            |
| Device classification | Catheter, Percutaneous (DQY)       |
| Date received         | Mar 2, 2000                        |
| Decision date         | Mar 31, 2000                       |
| Days to decision      | 29 days                            |
| Third-party review    | No                                 |
| Summary / Statement   | Summary                            |

**APPLICANT**

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|----------------|---|
| Company        | <b>Cordis Corp.</b>                                 |
| Location       | McHenry, IL, US                                     |
| Contact        | DENNIS S GRIFFIN                                    |
| Website        | <a href="https://cordis.com">https://cordis.com</a> |
| 510(k) history | 315 submissions · 281 cleared · 1976-2014           |

Cordis Corp. is a medical device manufacturer based in McHenry, US. The company specializes in interventional cardiovascular and gastroenterology devices. Cordis has a substantial FDA 510(k) regulatory history spanning from 1976 to 2014. The company received FDA 510(k) clearances from total submissions. Its portfolio focuses primarily on cardiovascular devices and gastroenterology stent systems, including percutaneous transluminal angioplasty catheters, emboli capture guidewires, and self-expanding biliary stent systems. Notable cleared products include the FLEXSTENT Bili...

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