

**K013963 MODEL 9210 DELIVERY CATHETER, MODEL 9210**Feb 28, 2002  
87 days to decisionK013963 · Product code: **DQY** · Cardiovascular  
Source: <https://www.510kdatabase.net/k013963/>**SUBMISSION DETAILS**

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|                       |                                    |
|-----------------------|------------------------------------|
| Decision              | Substantially Equivalent (Cleared) |
| Submission type       | Traditional                        |
| Device classification | Catheter, Percutaneous (DQY)       |
| Date received         | Dec 3, 2001                        |
| Decision date         | Feb 28, 2002                       |
| Days to decision      | 87 days                            |
| Third-party review    | No                                 |
| Summary / Statement   | Summary                            |

**APPLICANT**

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|----------------|---|
| Company        | <b>Medtronic Vascular</b>                 |
| Location       | Walker, MI, US                            |
| Contact        | MARY E BEST                               |
| 510(k) history | 475 submissions · 453 cleared · 1977-2023 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k013963/> 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 May 18, 2026