

**K925852 TERUMO CORONARY GUIDE WIRE**Jun 10, 1993  
205 days to decisionK925852 · Product code: **DQX** · CardiovascularSource: <https://www.510kdatabase.net/k925852/>**SUBMISSION DETAILS**

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|                       |                                    |
|-----------------------|------------------------------------|
| Decision              | Substantially Equivalent (Cleared) |
| Submission type       | Traditional                        |
| Device classification | Wire, Guide, Catheter (DQX)        |
| Date received         | Nov 17, 1992                       |
| Decision date         | Jun 10, 1993                       |
| Days to decision      | 205 days                           |
| Third-party review    | No                                 |
| Summary / Statement   | Summary                            |

**APPLICANT**

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|----------------|---|
| Company        | <b>Terumo Medical Corp.</b>               |
| Location       | Elkton, MD, US                            |
| Contact        | ALAN B HERSHMAN                           |
| 510(k) history | 143 submissions · 143 cleared · 1980-2011 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k925852/> 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 27, 2026