

**K260102 Tornier Perform Reversed Glenoid and Tornier Perform Reversed Augmented Glenoid Shoulder System**Jun 12, 2026  
150 days to decisionK260102 · Product code: **PHX** · Orthopedic  
Source: <https://www.510kdatabase.net/k260102/>**SUBMISSION DETAILS**

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|                       |  |
|-----------------------|--|
| Decision              | Substantially Equivalent (Cleared)               |
| Submission type       | Traditional                                      |
| Device classification | Shoulder Prosthesis, Reverse Configuration (PHX) |
| Date received         | Jan 13, 2026                                     |
| Decision date         | Jun 12, 2026                                     |
| Days to decision      | 150 days   |
| Third-party review    | No   |
| Summary / Statement   | Summary  |

**APPLICANT**

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|----------------|--|
| Company        | <b>Stryker Corporation (Tornier, Inc.)</b> |
| Location       | Bloomington, MN, US                        |
| Contact        | Jang Long                                  |
| 510(k) history | 3 submissions · 3 cleared · 2024-2026      |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k260102/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://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](https://accessdata.fda.gov). - Generated July 11, 2026