

**K920157 SCHNEIDER 8 FR. STAMINA(TM) GUIDING CATHETER**Mar 16, 1992  
62 days to decisionK920157 · Product code: **DXY** · CardiovascularSource: <https://www.510kdatabase.net/k920157/>**SUBMISSION DETAILS**

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|                       |                                             |
|-----------------------|---------------------------------------------|
| Decision              | Substantially Equivalent (Cleared)          |
| Submission type       | Traditional                                 |
| Device classification | Implantable Pacemaker Pulse-generator (DXY) |
| Date received         | Jan 14, 1992                                |
| Decision date         | Mar 16, 1992                                |
| Days to decision      | 62 days                                     |
| Third-party review    | No                                          |
| Summary / Statement   | Statement                                   |

**APPLICANT**

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|----------------|-----------------------------------------|
| Company        | <b>Schneider Intl., Ltd.</b>            |
| Location       | Minneapolis, MN, US                     |
| Contact        | ROBERT L ULLEN                          |
| 510(k) history | 22 submissions · 22 cleared · 1989-1995 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k920157/> 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 July 1, 2026