

**K953046 PERI-PUSHER**Sep 22, 1995  
84 days to decisionK953046 · Product code: **IYE** · Radiology  
Source: <https://www.510kdatabase.net/k953046/>**SUBMISSION DETAILS**

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|                       |                                    |
|-----------------------|------------------------------------|
| Decision              | Substantially Equivalent (Cleared) |
| Submission type       | Traditional                        |
| Device classification | Accelerator, Linear, Medical (IYE) |
| Date received         | Jun 30, 1995                       |
| Decision date         | Sep 22, 1995                       |
| Days to decision      | 84 days                            |
| Third-party review    | No                                 |
| Summary / Statement   | Statement                          |

**APPLICANT**

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|----------------|---|
| Company        | <b>Diacor, Inc.</b>                     |
| Location       | Salt Lake City, UT, US                  |
| Contact        | GLENN N WATERMAN                        |
| 510(k) history | 10 submissions · 10 cleared · 1986-2013 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k953046/> 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 17, 2026