

**K922942 PRO-SCOPE MODEL # PS-4000**Sep 10, 1993  
449 days to decisionK922942 · Product code: **EIA** · DentalSource: <https://www.510kdatabase.net/k922942/>**SUBMISSION DETAILS**

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|                       |                                    |
|-----------------------|------------------------------------|
| Decision              | Substantially Equivalent (Cleared) |
| Submission type       | Traditional                        |
| Device classification | Unit, Operative Dental (EIA)       |
| Date received         | Jun 18, 1992                       |
| Decision date         | Sep 10, 1993                       |
| Days to decision      | 449 days                           |
| Third-party review    | No                                 |
| Summary / Statement   | Statement                          |

**APPLICANT**

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|----------------|---------------------------------------|
| Company        | <b>Pro-Den Systems, Inc.</b>          |
| Location       | Portland, OR, US                      |
| Contact        | CADE ADAMS                            |
| 510(k) history | 2 submissions · 2 cleared · 1993-1995 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k922942/> 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 3, 2026