

**K033126 MONARCH 21 PENILE PLETHYSMOGRAPH SYSTEM**Oct 29, 2003  
29 days to decisionK033126 · Product code: **LIL** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k033126/>**SUBMISSION DETAILS**

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|                       |                                    |
|-----------------------|------------------------------------|
| Decision              | Substantially Equivalent (Cleared) |
| Submission type       | Special                            |
| Device classification | Monitor, Penile Tumescence (LIL)   |
| Date received         | Sep 30, 2003                       |
| Decision date         | Oct 29, 2003                       |
| Days to decision      | 29 days                            |
| Third-party review    | No                                 |
| Summary / Statement   | Summary                            |

**APPLICANT**

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|----------------|---------------------------------------|
| Company        | <b>Behavioral Technology, Inc.</b>    |
| Location       | Salt Lake City, UT, US                |
| Contact        | SUSAN E OLSEN                         |
| 510(k) history | 1 submissions · 1 cleared · 2003-2003 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k033126/> 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 June 30, 2026