

**K851646 VIBRECTOR**Aug 21, 1985  
124 days to decisionK851646 · Product code: **LIL** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k851646/>**SUBMISSION DETAILS**

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|-----------------------|------------------------------------|
| Decision              | Substantially Equivalent (Cleared) |
| Submission type       | Traditional                        |
| Device classification | Monitor, Penile Tumescence (LIL)   |
| Date received         | Apr 19, 1985                       |
| Decision date         | Aug 21, 1985                       |
| Days to decision      | 124 days                           |
| Third-party review    | No                                 |

**APPLICANT**

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|----------------|---------------------------------------|
| Company        | <b>Multicept, Aps.</b>                |
| Location       | Copenhagen, DK                        |
| Contact        | JAKOB WAGNER                          |
| 510(k) history | 2 submissions · 2 cleared · 1985-1996 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k851646/>; 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 29, 2026