

**K990125 HP VIRIDIA COMPONENT MONITORING SYSTEM,  
MODEL REV. K**Jan 29, 1999  
16 days to decisionK990125 · Product code: **DSI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k990125/>**SUBMISSION DETAILS**

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|                       |                                      |
|-----------------------|--------------------------------------|
| Decision              | Substantially Equivalent (Cleared)   |
| Submission type       | Special                              |
| Device classification | Detector And Alarm, Arrhythmia (DSI) |
| Date received         | Jan 13, 1999                         |
| Decision date         | Jan 29, 1999                         |
| Days to decision      | 16 days                              |
| Third-party review    | No                                   |
| Summary / Statement   | Summary                              |

**APPLICANT**

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|----------------|---|
| Company        | <b>Hewlett-Packard GmbH</b>             |
| Location       | 71004 Boblingen, DE                     |
| Contact        | EGON PFEIL                              |
| 510(k) history | 16 submissions · 16 cleared · 1995-2000 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k990125/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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