

**K991773 HP VIRIDIA COMPONENT MONITORING SYSTEM, HP VIRIDIA CMS 24/26, HP VIRIDIA MULTI-MEASUREMENT SERVER AND COMPACT PORTABLE P**Jun 7, 1999  
13 days to decisionK991773 · Product code: **DSI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k991773/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	May 25, 1999
Decision date	Jun 7, 1999
Days to decision	13 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/k991773/> 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 25, 2026