

**K980429 HP M2600A VIRIDIA TELEMETRY SYSTEM  
(FORMERLY KNOWN AS THE HP 2600A OMNI-CARE  
TELEMETRY SYSTEM)**Sep 9, 1998  
217 days to decisionK980429 · Product code: **DSI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k980429/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Feb 4, 1998
Decision date	Sep 9, 1998
Days to decision	217 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Hewlett-Packard Co.</b>
Location	Mchenry, IL, US
Contact	RAY STELTING
Website	<a href="https://www.hp.com">https://www.hp.com</a>
510(k) history	230 submissions · 229 cleared · 1976-2000

Hewlett-Packard Co. is a technology company headquartered in McHenry, US. The company historically developed medical devices alongside its core computing and printing business. Hewlett-Packard received FDA 510(k) clearances from total submissions, with clearances spanning 1976 to 2000. The company specialized in cardiovascular devices, including defibrillators, telemetry systems, and clinical information systems. Additional cleared devices covered gastroenterology, urology, and radiology applications. This regulatory record reflects the company's historical involvement in...