

**K992977 HEWLETT PACKARD MULTIFUNCTION PEDIATRIC  
DEFIB FUNCTION, MODEL M3503A & M3504A**Dec 1, 1999  
89 days to decisionK992977 · Product code: **MKJ** · Cardiovascular  
Source: <https://www.510kdatabase.net/k992977/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Sep 3, 1999
Decision date	Dec 1, 1999
Days to decision	89 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Hewlett-Packard Co.</b>
Location	Mchenry, IL, US
Contact	WARREN R WALTERS
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...

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Device record: <https://www.510kdatabase.net/k992977/>. Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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