

**K913451 HEWLETT-PACKARD MODEL 78720AC ARRHYTHMIA  
MONIT SYS**Oct 31, 1991  
90 days to decisionK913451 · Product code: **DSI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k913451/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Aug 2, 1991
Decision date	Oct 31, 1991
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Hewlett-Packard Co.</b>
Location	Mchenry, IL, US
Contact	WILLIAM C KOLE
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...