

**K864868 MICROPOROUS MONITORING ELECTRODES & ADAPTER BLOCK**Jan 14, 1987  
33 days to decisionK864868 · Product code: **DRX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k864868/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Dec 12, 1986
Decision date	Jan 14, 1987
Days to decision	33 days
Third-party review	No

**APPLICANT**

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Company	<b>Hewlett-Packard Co.</b>
Location	Mchenry, IL, US
Contact	DONALD A GUTHRIE
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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