

**K802718 MODEL 4700A ELECTROCARDIOGRAPH**Nov 26, 1980  
26 days to decisionK802718 · Product code: **DPS** · CardiovascularSource: <https://www.510kdatabase.net/k802718/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrocardiograph (DPS)
Date received	Oct 31, 1980
Decision date	Nov 26, 1980
Days to decision	26 days
Third-party review	No

**APPLICANT**

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Company	<b>Hewlett-Packard Co.</b>
Location	Mchenry, IL, US
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/k802718/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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