

**K861359 CARDIAC MONITOR, CARDIOTACHOMETER AND ALARM**May 16, 1986  
36 days to decisionK861359 · Product code: **DPS** · Cardiovascular  
Source: <https://www.510kdatabase.net/k861359/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrocardiograph (DPS)
Date received	Apr 10, 1986
Decision date	May 16, 1986
Days to decision	36 days
Third-party review	No

**APPLICANT**

---

Company	<b>Hewlett-Packard Co.</b>
Location	Mchenry, IL, US
Contact	NICHOLAS POLY
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...

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k861359/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 19, 2026