

**K822695 SIGNAL DISTRIBUTION NETWORK #78581A**Oct 15, 1982  
38 days to decisionK822695 · Product code: **DRT** · CardiovascularSource: <https://www.510kdatabase.net/k822695/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Sep 7, 1982
Decision date	Oct 15, 1982
Days to decision	38 days
Third-party review	No

**APPLICANT**

---

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...