

K952173 HEWLETT-PACKARD MODEL M1264B CATHSTATIONAug 11, 1995
95 days to decisionK952173 · Product code: **DPS** · CardiovascularSource: <https://www.510kdatabase.net/k952173/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrocardiograph (DPS)
Date received	May 8, 1995
Decision date	Aug 11, 1995
Days to decision	95 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Hewlett-Packard Co.
Location	McHenry, IL, US
Contact	ROB BUTLER
Website	https://www.hp.com
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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