FDA 510(k) Premarket Notification

CLEARED

K992636 HP CAREVUE 9000 CLINICAL INFORMATION SYSTEM, MODEL M2331A

Aug 31, 1999
25 days to decision

K992636 · Product code: **DXJ** · Cardiovascular
Source: https://www.510kdatabase.net/k992636/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Display, Cathode-ray Tube, Medical (DXJ)
Date received	Aug 6, 1999
Decision date	Aug 31, 1999
Days to decision	25 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Hewlett-Packard Co.**
Location	Mchenry, IL, US
Contact	MIKE HUDON
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...