FDA 510(k) Premarket Notification

K905788 MODEL M1251A MON. FULL DISCLOSURE REVIEW SYSTEM

CLEARED

Jun 6, 1991
161 days to decision

K905788 · Product code: **DXJ** · Cardiovascular
Source: https://www.510kdatabase.net/k905788/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Display, Cathode-ray Tube, Medical (DXJ)
Date received	Dec 27, 1990
Decision date	Jun 6, 1991
Days to decision	161 days
Third-party review	No

APPLICANT

Company	**Hewlett-Packard Co.**
Location	Mchenry, IL, US
Contact	RICHARD BEEBE
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...