K991871 HEWLETT PACKARD MULTI FUNCTION ADULT DEFIB ELECTRODE, MODEL M3501A & M3502A

CLEARED

Aug 5, 1999
65 days to decision

K991871 · Product code: **MKJ** · Cardiovascular
Source: https://www.510kdatabase.net/k991871/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Jun 1, 1999
Decision date	Aug 5, 1999
Days to decision	65 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Hewlett-Packard Co.**
Location	Mchenry, IL, US
Contact	WARREN R WALTERS
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...