FDA 510(k) Premarket Notification

K905101 NEONATAL/STERILE/DISPOS/NONINVAS BLOOD PRESS CUFF

CLEARED

Dec 20, 1990
37 days to decision

K905101 · Product code: **DXQ** · Cardiovascular
Source: https://www.510kdatabase.net/k905101/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Blood Pressure Cuff (DXQ)
Date received	Nov 13, 1990
Decision date	Dec 20, 1990
Days to decision	37 days
Third-party review	No

APPLICANT

Company	**Hewlett-Packard Co.**
Location	Mchenry, IL, US
Contact	DONALD A GUTHRIE
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...