FDA 510(k) Premarket Notification

K874421 PRE-GELLED DISPOSABLE ECG ELECTRODE - HP13941A

CLEARED

Jan 28, 1988
92 days to decision

K874421 · Product code: **DRX** · Cardiovascular
Source: https://www.510kdatabase.net/k874421/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Oct 28, 1987
Decision date	Jan 28, 1988
Days to decision	92 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...

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Device record: https://www.510kdatabase.net/k874421/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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