FDA 510(k) Premarket Notification

K820071 MODEL 5790 GAS CHROMATOGRAPH

CLEARED

Jan 28, 1982
16 days to decision

K820071 · Product code: **KZQ** · Toxicology
Source: https://www.510kdatabase.net/k820071/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Chromatography(gas), Clinical Use (KZQ)
Date received	Jan 12, 1982
Decision date	Jan 28, 1982
Days to decision	16 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k820071/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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