FDA 510(k) Premarket Notification

K761351 L080 SERIES LIQUID CHROMATOGRAPH

CLEARED

Feb 2, 1977
36 days to decision

K761351 · Product code: **KIE** · Toxicology
Source: https://www.510kdatabase.net/k761351/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Apparatus, High Pressure Liquid Chromatography (KIE)
Date received	Dec 28, 1976
Decision date	Feb 2, 1977
Days to decision	36 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...