FDA 510(k) Premarket Notification

K791893 19091-XXXXX SER. OF GLASS CAPILLARY COL.

CLEARED

Oct 30, 1979
35 days to decision

K791893 · Product code: **DII** · Toxicology
Source: https://www.510kdatabase.net/k791893/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Columns, Glc (DII)
Date received	Sep 25, 1979
Decision date	Oct 30, 1979
Days to decision	35 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...