

K781360 MODEL 80100A, PATIENT DATA MEMORYSep 7, 1978
31 days to decisionK781360 · Product code: **HGM** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k781360/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Monitoring, Perinatal (HGM)
Date received	Aug 7, 1978
Decision date	Sep 7, 1978
Days to decision	31 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...