

K921957 HEWLETT-PACKARD MODEL M1351A FETAL MONITORSep 24, 1993
515 days to decisionK921957 · Product code: **HGM** · Obstetrics & Gynecology
Source: <https://www.510kdatabase.net/k921957/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Monitoring, Perinatal (HGM)
Date received	Apr 27, 1992
Decision date	Sep 24, 1993
Days to decision	515 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Hewlett-Packard Co.
Location	Mchenry, IL, US
Contact	ERICH COURTIN
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...