

**K923343 HEWLETT-PACKARD MODEL M1020A PULSE
OXIMETER**Feb 10, 1993
237 days to decisionK923343 · Product code: **DQA** · Anesthesiology
Source: <https://www.510kdatabase.net/k923343/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Jun 18, 1992
Decision date	Feb 10, 1993
Days to decision	237 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Hewlett-Packard Co.
Location	Mchenry, IL, US
Contact	GERHARD LENKE
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...