FDA 510(k) Premarket Notification

K903523 HP MODEL M1020A, ARTERIAL OXYGEN SAT/PLETH MODULE

CLEARED

Oct 23, 1990
78 days to decision

K903523 · Product code: **DQA** · Anesthesiology
Source: https://www.510kdatabase.net/k903523/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Aug 6, 1990
Decision date	Oct 23, 1990
Days to decision	78 days
Third-party review	No

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...

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k903523/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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