

**K881136 HEWLETT-PACKARD MODELS 78354A & 78352A
PULSE OXI.**May 18, 1988
63 days to decisionK881136 · Product code: **DQA** · Anesthesiology
Source: <https://www.510kdatabase.net/k881136/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Mar 16, 1988
Decision date	May 18, 1988
Days to decision	63 days
Third-party review	No

APPLICANT

Company	Hewlett-Packard Co.
Location	McHenry, IL, US
Contact	NICHOLAS POLY
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...

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