FDA 510(k) Premarket Notification

K883606 MODEL HP 15209A TRANS. COMBINED TCP02/TCPCO2 TRANS

CLEARED

Dec 16, 1988
115 days to decision

K883606 · Product code: **LKD** · Anesthesiology
Source: https://www.510kdatabase.net/k883606/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Carbon-dioxide, Cutaneous (LKD)
Date received	Aug 23, 1988
Decision date	Dec 16, 1988
Days to decision	115 days
Third-party review	No

APPLICANT

Company	**Hewlett-Packard Co.**
Location	Mchenry, IL, US
Contact	DONALD J BARTH
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...