

**K883481 HP 77025A/77030A ULTRA. SYSTEM/HP 77020CV  
UPGRADE**Nov 4, 1988  
80 days to decisionK883481 · Product code: **IYO** · Radiology  
Source: <https://www.510kdatabase.net/k883481/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Echo, Ultrasonic (IYO)
Date received	Aug 16, 1988
Decision date	Nov 4, 1988
Days to decision	80 days
Third-party review	No

**APPLICANT**

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Company	<b>Hewlett-Packard Co.</b>
Location	Mchenry, IL, US
Contact	CHARLES R BURR
Website	<a href="https://www.hp.com">https://www.hp.com</a>
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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