

**K895026 HP MODEL M1318A TRANSVAGINAL ULTRASOUND
TRANSDUCER**Nov 6, 1989
90 days to decisionK895026 · Product code: **HGL** · Obstetrics & Gynecology
Source: <https://www.510kdatabase.net/k895026/>**SUBMISSION DETAILS**

| | |
|-----------------------|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Transducer, Ultrasonic, Obstetric (HGL) |
| Date received | Aug 8, 1989 |
| Decision date | Nov 6, 1989 |
| Days to decision | 90 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---|
| Company | Hewlett-Packard Co. |
| Location | Mchenry, IL, US |
| Contact | CHARLES R BURR |
| 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...
