

**K970456 HEWLETT APCKARD SERIES 50 OB TRACEVUE  
OBSTETRICAL SURVEILLANCE AND ARCHIVING SYSTEM**Oct 3, 1997  
239 days to decisionK970456 · Product code: **HGM** · Obstetrics & Gynecology  
Source: <https://www.510kdatabase.net/k970456/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Monitoring, Perinatal (HGM)
Date received	Feb 6, 1997
Decision date	Oct 3, 1997
Days to decision	239 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Hewlett-Packard GmbH</b>
Location	71004 Boblingen, DE
Contact	MIKE HUDON
510(k) history	16 submissions · 16 cleared · 1995-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k970456/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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