FDA 510(k) Premarket Notification

CLEARED

K970456 HEWLETT APCKARD SERIES 50 OB TRACEVUE OBSTETRICAL SURVEILLANCE AND ARCHIVING SYSTEM

Oct 3, 1997
239 days to decision

K970456 · Product code: **HGM** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k970456/

SUBMISSION DETAILS

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

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