FDA 510(k) Premarket Notification

K990125 HP VIRIDIA COMPONENT MONITORING SYSTEM, MODEL REV. K

CLEARED

Jan 29, 1999
16 days to decision

K990125 · Product code: **DSI** · Cardiovascular
Source: https://www.510kdatabase.net/k990125/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Jan 13, 1999
Decision date	Jan 29, 1999
Days to decision	16 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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