FDA 510(k) Premarket Notification

K971910 MULTI-PARAMETER MODULE (M3000A)/DISPLAY UNIT (M3046A)

CLEARED

Mar 23, 1998
304 days to decision

K971910 · Product code: **DSI** · Cardiovascular
Source: https://www.510kdatabase.net/k971910/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	May 23, 1997
Decision date	Mar 23, 1998
Days to decision	304 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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