

**K971910 MULTI-PARAMETER MODULE (M3000A)/DISPLAY  
UNIT (M3046A)**Mar 23, 1998  
304 days to decisionK971910 · Product code: **DSI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k971910/>**SUBMISSION DETAILS**

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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**

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Company	<b>Hewlett-Packard GmbH</b>
Location	71004 Boblingen, DE
Contact	EGON PFEIL
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/k971910/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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