

K001722 AGILENT COMPONENT MONITORING SYSTEMJun 30, 2000
24 days to decisionK001722 · Product code: **MHX** · CardiovascularSource: <https://www.510kdatabase.net/k001722/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Monitor, Physiological, Patient(with Arrhythmia Detection Or Alarms) (MHX)
Date received	Jun 6, 2000
Decision date	Jun 30, 2000
Days to decision	24 days
Third-party review	No
Summary / Statement	Summary
Other names	AGILENT INVASIVE BLOOD PRESSURE MODULE

APPLICANT

Company	Agilent Technologies Deutschland GmbH
Location	Boeblingen,, DE
Contact	EGON PFEIL
510(k) history	4 submissions · 4 cleared · 2000-2001

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k001722/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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