

**K003038 AGILENT COMPONENT MONITORING SYSTEM,
AGILENT V24/26, AGILENT BIS MODULE, MODEL
M1175A/76A/77A,REV.L,M1205A,REV.L,M1034A**

Dec 15, 2000
77 days to decision

K003038 · Product code: **OLW** · Neurology
Source: <https://www.510kdatabase.net/k003038/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Index-generating Electroencephalograph Software (OLW)
Date received	Sep 29, 2000
Decision date	Dec 15, 2000
Days to decision	77 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k003038/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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