

Audit Report: 21 CFR Part 11

tiamo 2.4



Metrohm AG
CH-9100 Herisau
Switzerland
Tel. +41 71 353 85 85
Fax +41 71 353 89 01
www.metrohm.com

Audit Date:	05.09.2013 ¹
Auditor:	Sieghard Wagner, mech. Engineer (grad.), Chemengineering Business Design GmbH
Audit Target:	Compliance check of <i>tiamo 2.4</i> software for controlling titration devices against the requirements of 21 CFR Part 11.
Description:	PC software for controlling titration devices. Creation of methods, carrying out determinations (titrations and measurements), data acquisition, evaluation and archiving in a database. The above software was developed and manufactured in accordance with the requirements demanded by the ISO 9001 quality system regarding the design, manufacture and servicing of Metrohm instruments.
Investigation:	The functions and properties of the above software system are audited with the requirements of 21 CFR Part 11 and the current interpretations. During the audit the following items were examined: Validation, password, documentation, audit trail, electronic copies, access rights, user access, access violation, sequence of steps, plausibility, device check, data encryption, electronic signature.
Operator Responsibility:	The compliance with the requirements of 21 CFR Part 11 can only be met together with the operational environment. The requirements for such an environment are: - technical environment, data management - training - administration - SOP (Standard Operating Procedures)
Summary:	The software is compliant to the following 21 CFR Part 11 requirements: 11.10 (b), (d), (e), (f), (g), (h) 11.50, 11.70 11.300 (a). The software is compliant to the following 21 CFR Part 11 requirements with support of the operator. 11.10 (a), (c), (i), (j), (k) 11.100 (a), (b), 11.200 (a), 11.300 (b), (c), (d). The software is supporting electronic signature.

Sieghard Wagner

¹ This 21 CFR Part 11 certificate is based on a physical audit performed January the 13th 2009. According to Metrohm AG, the implemented changes in the current version are not relevant with regard to 21 CFR Part 11 or compliant with 21 CFR Part 11 (see Release Notes 8.101.8017EN, 8.101.8027EN, 8.101.8039EN, 8.101.8055EN). Therefore, this update does not require a physical re-audit.