

Audit Report: 21 CFR Part 11

tiamo 2.5



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Audit Date:	07.07.2015 ¹
Auditor:	Sieghard Wagner, mech. Engineer (grad.), Chemgineering Business Design GmbH
Audit Target:	Proof that the <i>tiamo 2.5</i> software, used for control of titration devices, is in compliance with 21 CFR Part 11.
Description:	<p><i>tiamo</i> is a PC software for control of titration devices. It features creating methods, performing determinations (titrations and measurements), data acquisition, evaluation and archiving in a database.</p> <p>The software was developed by Metrohm AG in accordance with ISO 9001 requirements regarding design, manufacturing and maintenance.</p>
Investigation:	<p>The functions and properties of the above software system are audited with the requirements of 21 CFR Part 11 and the current interpretations.</p> <p>During the audit the following items were examined:</p> <ul style="list-style-type: none">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:	<p>Compliance with the requirements of 21 CFR Part 11 can only be met in connection with the operational environment. The requirements for such an environment encompass:</p> <ul style="list-style-type: none">technical environment, data managementtrainingadministrationStandard Operating Procedures (SOP)
Summary:	<p>The software is compliant with the following 21 CFR Part 11 requirements:</p> <ul style="list-style-type: none">11.10 (b), (d), (e), (f), (g)11.50, 11.7011.300 (a). <p>The software is compliant with the following 21 CFR Part 11 requirements with support of the operator.</p> <ul style="list-style-type: none">11.10 (a), (c), (h), (i), (j), (k)11.100 (a), (b), 11.200 (a), 11.300 (b), (c), (d). <p>The software supports electronic signatures.</p>

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, 8.101.8072EN). Therefore, this update does not require a physical re-audit.