



Security Software for NanoDrop One Spectrophotometers

Enables Title 21 CFR Part 11 compliance

PC Control software for NanoDrop One/One^c spectrophotometers enables 21 CFR Part 11 compliance

NanoDrop One PC Control software allows analysts to operate the spectrophotometer from a computer as well as the built-in touch screen. Security Suite software, a proven compliance component of many other Thermo Scientific systems, integrates directly into the NanoDrop One PC Control software so analysts can confidently operate the trusted NanoDrop One/One^c spectrophotometer in compliance with regulatory requirements.

Security Suite for NanoDrop One/One^c Spectrophotometers

Security Suite software for NanoDrop One/One^c Spectrophotometers can integrate into new and older instruments so all users can be compliant with this new software.

➤ To purchase the Security Suite software, **please contact us** or visit witec.ch/Nanodrop

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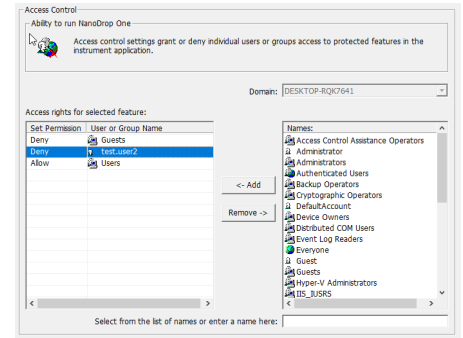
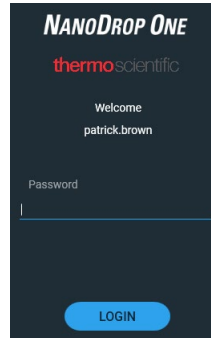
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Security Suite for the NanoDrop One/One^c PC Control software makes compliance easy

Compliance doesn't have to be complicated. The table below shows an example of how Security Suite manages regulatory requirements.

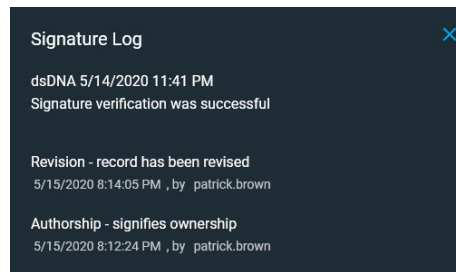
“Limiting system access to authorized individuals.”



Administrators grant access through the Security Suite software. User account access is managed through the Windows login system.

“Signed electronic records shall contain information associated with the signing that clearly indicates all of the following:

1. The printed name of the signer;
2. The date and time when the signature was executed; and
3. The meaning (such as review, approval, responsibility, or authorship) associated with the signature.”

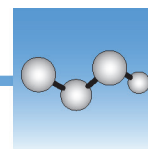


Experiments are signed immediately after they are completed. Users can also revise, review, and approve experiments from the History page.

“Use of secure, computer-generated, time-stamped audit trails to independently record the date and time of operator entries and actions that create, modify, or delete electronic records. Record changes shall not obscure previously recorded information. Such audit trail documentation shall be retained for a period at least as long as that required for the subject electronic records and shall be available for agency review and copying.”

Audit Event Report					15-May-2020T23:49:48+08:00
Source	User Name	Severity	Timestamp Local +08:00	Category	
Audit Manager	patrick.brown	Information	15-May-2020T23:49:14	Signature	
Description	The file "Reports at 15-May-2020 23:49:10 (GMT+08:00)" was signed. Reason : Authorship - signifies ownership				
NanoDrop One	patrick.brown	Information	15-May-2020T20:14:06	Signature	
Description	The record "Experiment dsDNA 5/14/2020 11:41 PM." was signed. Reason : Revision - record has been revised				
NanoDrop One	patrick.brown	Information	15-May-2020T20:13:29	Signature	
Description	The record "Experiment NucleicAcid_5/15/2020 5:04 AM." was signed. Reason : Approval - record is approved for use				

The Audit Manager creates an electronic audit trail of actions such as logons, data creation, signatures, and more. Audit logs reports can be generated and printed as needed.



21 CFR Part 11 Compliance

This document explains how the following Thermo Scientific™ software can help you comply with the regulations in 21 CFR Part 11¹ for electronic records and electronic signatures.

Table 1. Thermo Scientific software applications that offer 21 CFR Part 11 compliance

Application Name	Application Name
OMNIC™ Paradigm	OMNIC™xi
OMNIC™	TQ ANALYST™
OMNIC™ for DISPERSIVE RAMAN™	RESULT™
OMNIC™ SPECTA™	INSIGHT™
OMNIC™ PICTA™	CUE™
NanoDrop One™	

History

Part 11 of the 21 CFR (Title 21 – Food and Drugs of the Code of Federal Regulations) is a document issued by the United States Food and Drug Administration (FDA) that outlines the FDA criteria for accepting electronic records and signatures. The regulations in the final version of 21 CFR Part 11 became effective on August 20, 1997. All industries, companies and organizations regulated by the FDA that utilize electronic records must follow these regulations.

In 1991 the FDA met with representatives from the pharmaceutical industry to determine how to accommodate an electronic record system, under the guidelines of current Good Manufacturing Practice (cGMP), that would create a “paperless” record system. The primary concerns of the FDA were maintaining the trustworthiness, reliability, and integrity of the electronic records and ensuring that electronic records were equivalent to paper records. The 21 CFR Part 11 regulation was created to prevent fraud in the generation and signing of electronic records.

¹ 21 CFR Part 11 Electronic Records; Electronic Signatures; Final Rule,” Federal Register 62, no. 54 (1997): 13430-13556. World Wide Web <http://www.fda.gov>.

Definitions

Understanding the following terms is essential for the successful implementation of the regulations in 21 CFR Part 11. These definitions taken directly from 21 CFR Part 11 will be the starting point for our discussion of Thermo Scientific's software compliance with the regulation.

Closed system—An environment in which system access is controlled by persons who are responsible for the content of electronic records that are on the system.

Open system—An environment in which system access is not controlled by persons who are responsible for the content of electronic records that are on the system.

Digital signature (DS)—Electronic signature based upon cryptographic methods of originator authentication, computed by using a set of rules and a set of parameters such that the identity of the signer and the integrity of the data can be verified.

Electronic record—Any combination of text, graphics, data, audio, pictorial, or other information representation in digital form that is created, modified, maintained, archived, retrieved, or distributed by a computer system.

Electronic signature—A computer data compilation of any symbol or series of symbols executed, adopted, or authorized by an individual to be the legally binding equivalent of the individual's handwritten signature.

Software application template files—May include parameter files, quant methods and macros.

Key subparts of Part 11: Electronic records; Electronic signatures

- 21 CFR Part 11 is divided into three subparts.
- Subpart A defines the scope and implementation of the regulations and defines key terms in the document.
- Requirements for electronic records are described in Subpart B, including controls for data generation from closed and open systems as well as information associated with the electronic signature and linking the signature to the record.
- Subpart C details the requirements, components and controls of electronic signatures.

Software

Every Thermo Scientific security software package is designed under the strict guidelines of Thermo Fisher Scientific's ISO 9001 certified product development process at our development and manufacturing site in Madison, Wisconsin. Trained members from different functional departments at our facility adhere to quality guidelines covering all aspects of

development. Each software development project begins with specifications created with our customers' needs in mind. The software designs are based on object-oriented and modular architecture. Software development practices follow our Product Development Process, which includes procedures for change control, source-code control systems, and defect management. Complete user documentation is created for every project. Intensive verification and regression testing of the software is performed according to the project test plan. The qualification package can be used to verify the consistency and accuracy of the spectrophotometer's operation compared with specified limits.

21 CFR Part 11 Compliance Statement

When the software application is installed, the following tools will help you achieve compliance with 21 CFR Part 11 in a laboratory setting:

- System log-ins and passwords
- Complete access control over software features in an easy-to-use interface
- An extensive set of software policies that allow control over program and file or data record operations
- Embedded spectral history tracking, including user information, spectrophotometer parameters, and any data manipulation information produced throughout the life of the file or data record
- Complete software use and event audit trails using a custom log, even when the software is not running
- Digital signatures with reports, data, macros, configurations, TQ Analyst quantitative and qualitative methods, and experiment files

Note Macros, configurations and experiment files do not apply to the OMNIC Paradigm, INSIGHT, RESULT, or OMNIC Spectra software applications.

- Network and local security with Windows® software
- Data record overwrite protection
- The ability to detect changes or data tampering through encrypted digital signatures

Windows security is embedded in the software structure and is set up through the Windows security features. You can control access to an instrument by using those features in conjunction with software access privileges. The Windows login and password are used to authenticate users when an electronic record is created. Those responsible for maintaining system records must take measures to ensure the software operates in a closed system.

The following sections explain how you can use the software tools listed above to help you meet each requirement of the 21 CFR Part 11 regulation. Certain sections of the regulation are solely the responsibility of the owner of the system; we cannot directly provide tools for compliance with those specific sections. It is important to note that compliance with 21 CFR Part 11 extends beyond software implementation and will require laboratory and computer procedures that control all phases of electronic record creation and management.

Note In this document, specific requirements from the 21 CFR Part 11 Electronic Records/Electronic Signature rule are shown in italics with quotation marks. Our capability to meet these requirements is shown in plain text after the statement of the requirement.

Part 11: Electronic records; Electronic signatures

This section covers Subparts B and C of 21 CFR Part 11.

Subpart B: Electronic records

§11.10 Controls for closed systems

Persons who use closed systems to create, modify, maintain, or transmit electronic records shall employ procedures and controls designed to ensure the authenticity, integrity, and, when appropriate, the confidentiality of electronic records, and to ensure that the signer cannot repudiate the signed record as not genuine. Such procedures and controls shall include the following:

(a) “Validation of systems to ensure accuracy, reliability, consistent intended performance, and the ability to discern invalid or altered records.”

The system owner must develop a protocol for validating the system. We offer a suite of products and services for the qualification of your laboratory. These offerings provide you with the tools, documentation and certification services that make system qualification efforts progress smoothly.

The ability to detect invalid or altered records is controlled by using the digital signature option in the software. With digital signatures, result data, application template files and qualification test files can be digitally signed, ensuring the validity of the record. By checking for the presence of a correct digital signature, the software can detect invalid or altered records.

(b) “The ability to generate accurate and complete copies of records in both human readable and electronic form suitable for inspection, review, and copying by the agency. Persons should contact the agency if there are any questions regarding the ability of the agency to perform such review and copying of the electronic record.”

When data is collected, detailed information about the experiment, instrument and accessory is stored in a non-editable, file-embedded spectral history. If and when the data is post-processed in any manner, details about the processing operation are noted in the data's history. Information about system user and digital signatures is also stored. You can view and print the data files and their history at any time, if desired. The system owner is responsible for which format will be used to save data.

OMNIC Paradigm software also provides a record of detailed information to be stored in a database. The database includes all data record information listed above, and data records and history can be viewed and printed at any time using the Audit Manager software.

(c) "Protection of records to enable their accurate and ready retrieval throughout the records retention period."

You can store data created by the software directly to a secure server. The system owner or IT group must determine how the files will be archived and who has access to these records. OMNIC Paradigm software offers data storage directly to a database on a secure server. All other applications offer secure data file storage directly to a secure server.

(d) "Limiting access to authorized individuals."

Using Windows secure logins are required for controlling access to the system. The software must be installed on computers that have the supported Windows operating system, Windows users and groups can be configured to have access to the software through the Security Administration program.

The login feature of Windows software allows system administrators to restrict system access to only authorized users. To gain access to the software, users must log into Windows software with their user name and password. The Windows password must be reentered when launching the software. To ensure full security, the user shall be given a unique user name and a private password.

The system administrator can configure users' profiles to restrict their software access to only the programs they need. A Windows system must be configured with a secure file system in order to grant individual read, write and delete access to users. The Security Administration software is used to set access privileges to data records, features, security policies and signature meanings. Control of file operations on the computer that are conducted external to the software application is the responsibility of the system owner.

(e) "Use of secure, computer-generated, time-stamped audit trails to independently record the date and time of operator entries and action that create, modify, or delete electronic records. Record changes shall not obscure previously recorded information. Such audit trail documentation shall be retained for a period at least as long as that required for the subject electronic records and shall be available for agency review and copying."

When data is collected, detailed information about the date and time, operator, experiment, instrument and accessory are stored in a non-editable, file- or database-embedded spectral history. (In addition to the spectral history, OMNIC Paradigm software also stores this information in its database.) Test files are used to perform instrument qualification. These test files contain the sequence of events for the qualification and are signed at the factory.

The history provides an internal recording of all data manipulations for any given data file after it is created. In addition, the Thermo Scientific Audit Log Service logs all Thermo Fisher Scientific (TFS) file operations, both within and outside of the software applications. Thus, the Security Suite will log any attempt to create, modify or delete any TFS records on the system.

(f) "Use of operational system checks to enforce permitted sequencing of steps and events, as appropriate."

The software can enforce step sequencing and events for all aspects of data collection, processing and archiving. Sequenced step files can be created in the software. These files can specify and sequence the complete process: collection of parameters, data collection, final formats, post-processing operations, and archiving of data. The software can be configured so that users can access only specific files.

The Windows user access privileges are extended into the application and Security Administration software. Membership in the system group can control the extent of software access of each user. The system administrator or IT group must establish access failure criteria, and it is the responsibility of the Windows administrator to set these security features to ensure only authorized individuals have access to the system. If the sequence in a test file has been altered, the software notifies the user, the signature is invalidated, and the test will not run.

(g) "Use of authority checks to ensure that only authorized individuals can use the system, electronically sign a record, access the operation or computer system input or output device, alter a record, or perform the operation at hand."

The system administrator or IT group must establish access failure criteria, and it is the responsibility of the Windows administrator to set these security features to ensure that only authorized individuals have access to the system and files on the system.

(h) "Use of device (e.g., terminal) checks to determine, as appropriate, the validity of data input or operational instruction."

The system firmware and software application template files determine the validity of the input or operational instruction of the instrument. The system firmware is an IQ-qualified standard component of the instrument and can only be updated by a trained and certified Thermo Scientific service representative. Changes to the software application template files are controlled by user-access policies set up by the system administrator. Administrators can save the software application template files in a secure Windows file system to prevent unauthorized users from changing the operation parameters of the system.

(i) “Determination that persons who develop, maintain, or use electronic record/electronic signature systems have the education, training, and experience to perform their assigned tasks.”

We train our system developers and maintain training records according to our internal training procedure. A training matrix is maintained along with individual training records for each developer. Thermo Fisher Scientific is ISO 9001 certified and follows these guidelines when developing all products.

Our service representatives must be trained in order to maintain and service our instruments and software. Service representatives receive training on the qualification and security software and must be recertified every two years. A training matrix is also maintained for our service representatives.

It is the responsibility of the system owner to determine that persons who develop, maintain, or use electronic record/electronic signature systems at your site have the education, training and experience to perform their assigned task.

(j) “The establishment of, and adherence to, written policies that hold individuals accountable and responsible for actions initiated under their electronic signatures, in order to deter record and signature falsification.”

To deter falsification or fraud, the system owner must establish written policies that hold individuals accountable for actions initiated under electronic signatures.

(k1) Use of appropriate controls over systems documentation including: Adequate controls over the distribution of, access to, and use of documentation for system operation and maintenance.”

The software is supplied with documentation for the operation and maintenance of the our instruments and software. You can use the information in the documentation to create Standard Operating Procedures (SOPs). It is the responsibility of the system owner to control the system documentation.

(k2) Use of appropriate controls over systems documentation including: Revision and change control procedures to maintain an audit trail that documents time-sequenced development and modification of systems documentation.”

Our documentation contain version information that can be incorporated into the system owner’s documentation control system. You can obtain information about software and firmware version numbers by choosing “About” in the Help menu. The system owner must implement a change control protocol for system documentation.

§11.30 Controls for open systems

“Persons who use open systems to create, modify, maintain, or transmit electronic records shall employ procedures and controls designed to ensure the authenticity, integrity, and, as appropriate, the confidentiality of electronic records from the point of their creation to the point of their receipt. Such procedures and controls shall include those identified in § 11.10, as appropriate, and additional measures such as document encryption and use of appropriate digital signature standards to ensure, as necessary under the circumstances, record authenticity, integrity, and confidentiality.”

The software implementation requires the use of a closed system since we do not employ data encryption on our data files. Windows security is embedded in the software structure, and security is set up through the Windows security feature. The Windows login and password, in conjunction with the password reverification required when a user starts the software, provide a way to control access to the software and an instrument. By following the guidelines in this document, you can achieve compliance with 21 CFR Part 11 as it pertains to a closed system.

Although data encryption is not used, the system administrator may choose to store the data on a secure server (recommended) such that only authorized users may access data according to their privileges. These privileges must be controlled by a unique user name and password combination.

If compliance is desired in an open system, those responsible for maintaining system records must take adequate measures to ensure that the software complies.

§11.50 Signature manifestations

(a) “Signed electronic records shall contain information associated with the signing that clearly indicates all of the following:

- (1) The printed name of the signer;
- (2) The date and time when the signature was executed; and
- (3) The meaning (such as review, approval, responsibility, or authorship) associated with the signature.”

All digital signatures produced by the software contain the information specified by the regulations, in addition to the signature.

(b) “The items identified in paragraphs (a)(1), (a)(2), and (a)(3) of this section shall be subject to the same controls as for electronic records and shall be included as part of any human readable form of the electronic record (such as electronic display or printout).”

Since digital signatures implemented in the software are embedded within the electronic record, these signatures are subject to the same controls as the electronic record. The signature is included as part of the human readable and printed form of the electronic record.

§11.70 Signature/Record Linking

“Electronic signatures and handwritten signatures executed to electronic records shall be linked to their respective electronic records to ensure that the signatures cannot be excised, copied, or otherwise transferred to falsify an electronic record by ordinary means.”

The digital signature is stored in the same data file or report that is signed. Because the signature is stored in the same file as the electronic record, all digital signatures produced in the software are directly linked to the electronic record. A check of the electronic record can reveal an invalid electronic record.

Subpart C: Electronic signatures

§11.100 General Requirements for Electronic Signatures

(a) “Each electronic signature shall be unique to one individual and shall not be reused, or reassigned, to anyone else.”

The system owner’s policy for assigning Windows user IDs and passwords must comply with this requirement, which can be accomplished by assigning a unique user name to each individual and by not reusing or reassigning any user names. If the user names are unique for all individuals with access to the system, the digital signature produced by the software will be unique.

(b) “Before an organization establishes, assigns, certifies, or otherwise sanctions an individual’s electronic signature, or any element of such signature, the organization shall verify the identity of the individual.”

The system owner must take appropriate measures to ensure the identity of all individuals who may be involved in applying electronic signatures to records.

(c) “Persons using electronic signatures shall, prior to or at the time of such use, certify to the agency that the electronic signatures in their system, used on or after August 20, 1997, are intended to be the legally binding equivalent of traditional handwritten signatures.

(1) The certification shall be submitted in paper form and signed with a traditional handwritten signature, to the Office of Regional Operations (HFC–100), 5600 Fishers Lane, Rockville, MD 20857.

(2) Persons using electronic signatures shall, upon agency request, provide additional certification or testimony that a specific electronic signature is the legally binding equivalent of the signer’s handwritten signature.”

In order to have an electronic signature, the organization using the signature must make it legally binding by submitting a letter and a form to the FDA.

§11.200 Electronic Signature Components and Controls

(a) “Electronic signatures not based upon biometrics shall:

(1) Employ at least two distinct identification components such as an identification code and password.

(i) When an individual executes a series of signings during a single, continuous period of controlled system access, the first signing shall be executed using all electronic signature components; subsequent signings shall be executed using at least one electronic signature component that is only executable by, and designed to be used only by, the individual.

(ii) When an individual executes one or more signings not performed during a single, continuous period of controlled system access, each signing shall be executed using all of the electronic signature components.

(2) Be used only by their genuine owner; and

(3) Be administered and executed to ensure that attempted use of an individual's electronic signature by anyone other than its genuine owner requires collaboration of two or more individuals."

Digital signatures used by the software are based on the user's log in ID and Windows password. Windows software is used to generate the digital signature in the software, and the combination of the signature components is unique for each user, as long as the requirements in 11.100 (a) are met. All signings in the software require entering the password of the person who is logged in to the Windows session at the time of system use. The system owner and administration must implement a protocol for using electronic signatures as described in requirements (2) and (3) are met.

(b) "Electronic signatures based on biometrics shall be designed to ensure that they cannot be used by anyone other than their genuine owners."

This requirement does not apply to our software because we use digital signatures based on the combination of a user name and password, instead of biometrics.

\$11.300 Controls for identification codes/passwords

"Persons who use electronic signatures based upon use of identification codes in combination with passwords shall employ controls to ensure their security and integrity. Such controls shall include:

(a) "Maintaining the uniqueness of each combined identification code and password, such that no two individuals have the same combination of identification code and password."

The system administrator or IT group must ensure that the combination of ID code and password is unique for each individual. This can be easily accomplished by issuing each user a unique login identification.

(b) "Ensuring that identification code and password issuances are periodically checked, recalled, or revised (e.g., to cover such events as password aging)."

(c) "Following loss management procedures to electronically deauthorize lost, stolen, missing, or otherwise potentially compromised tokens, cards, and other devices that bear or generate identification code or password information, and to issue temporary or permanent replacements using suitable, rigorous controls."

(d) "Use of transaction safeguards to prevent unauthorized use of passwords and/or identification codes, and to detect and report in an immediate and urgent manner any attempts at their unauthorized use to the system security unit, and, as appropriate, to organizational management."

Windows security features simplify the process of periodic checking, recalling and revising of log-ins and passwords. Transaction safeguards to prevent unauthorized access to the system are also available in the Windows operating system.

Limiting the number of failed login attempts and creating a password for an aging procedure is a common safeguard to limit and log the number of failed login attempts. Consult your Windows documentation for more information about activating system safeguards. The system administrator must establish a procedure for checking ID codes and passwords and loss management.

(e) “Initial and periodic testing of devices, such as tokens or cards, that bear or generate identification code or password information to ensure that they function properly and have not been altered in an unauthorized manner.”

Because our software does not use cards or tokens to generate identification codes, this requirement does not apply.

Summary

This document was created based on Thermo Fisher Scientific’s interpretation of the regulations and through consultation with experts in the field. The software with the digital signature option can be used together with proper procedures and controls instituted by our customers, in accordance with an FDA compliant process.

More About 21 CFR Part 11

For more information about the requirements of 21 CFR Part 11, go to www.fda.gov.

We are dedicated to working with our customers to help meet their regulatory needs wherever possible. For more information contact: www.thermofisher.com

21 CFR Part 11 Compliance

More About 21 CFR Part 11

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