



Sparta Systems TrackWise Solution

21 CFR Part 11 and Annex 11 Assessment

October 2017

Introduction

The purpose of this document is to outline the roles and responsibilities for compliance with the FDA's 21 CFR Part 11 and the European Union's Annex 11 as they apply to Sparta System's TrackWise product. The regulations require organizations to have administrative, procedural and technical controls in place. While it is not possible for Sparta to offer a turnkey 21 CFR Part 11 or EU Annex 11 compliant system, the recommendations in this document will assist using organizations in achieving compliance.

Both regulations cover the same topic, the use of computerized systems in regulatory environments. However, the approach of 21 CFR Part 11 is to clarify the requirements to be met with an emphasis on activities and reporting. EU Annex 11 points to risk assessment as the start of compliance activities. In addition, Part 11 differentiates security for open and closed systems, with security for open systems but without reference to risk and criticalities. The aggregate of these differences is represented with the comparison matrix shown below.

High-level Comparison of EU Annex 11 and FDA 21 CFR Part 11

	Part 11	Annex 11
Scope/Principle	Electronic records and electronic signatures as used for all FDA regulated activities.	Computerized systems as part of GMP regulated activities. Application should be validated. IT infrastructure should be qualified.
Focus	Using electronic records and signatures in open and closed computer systems.	Risk- based quality management of computerized systems.
Objective	Electronic records and signatures should be as trustworthy and reliable as paper records and handwritten signatures.	Using a computerized system should ensure the same product quality and quality assurance as manual systems with no increase in the overall risk.

Procedures and Controls for Closed Systems

21 CFR Part 11	Annex 11	Responsible Party	TrackWise
<p>11.10(a)</p> <p>Is the system validated?</p>	<p>4.1</p> <p>Do validation documents and reports cover the relevant steps of the life cycle?</p> <p>4.2</p> <p>Do validation documents include change control records (if applicable) and reports on deviations observed during the validation process?</p>	User	<p>Validation is the overall responsibility of the using organization.</p> <p>Sparta Systems, Inc. does its own internal testing and validation before each release in accordance with documented SOPs.</p> <p>This validation covers the core usage of the system with baseline configuration; customers must validate any additional configuration created.</p>
<p>11.10(a)</p> <p>Is it possible to discern invalid or altered records?</p>		Sparta	<p>TrackWise offers a full audit trail where relevant changes are logged. The audit trail includes user ID, old and new value and time stamp. Unauthorized changes are prevented by the access security controls.</p>
<p>11.10(b)</p> <p>Is the system capable of producing accurate and complete copies of electronic records on paper?</p>	<p>8.1</p> <p>Is the system capable of producing clear printed copies of electronically stored data?</p>	Sparta	<p>Full reporting capability that can be printed on paper or produced electronically.</p>
<p>11.10(b)</p> <p>Is the system capable of producing accurate and complete copies of records in electronic form for inspection, review, and copying by the FDA?</p>		Sparta	<p>Records can be saved electronically in Rich Text Format (rtf); Adobe Portable Document Format (pdf); MS Word (doc); MS Excel (xls); and Crystal Reports (rpt) format.</p>

21 CFR Part 11	Annex 11	Responsible Party	TrackWise
<p>11.10(c)</p> <p>Are the records readily retrievable throughout their retention period?</p>	<p>17</p> <p>Is data archived? If data is archived is it checked for accessibility, readability and integrity? When changes are made to the system, is the ability to retrieve archived data ensured and tested?</p>	<p>User & Sparta</p>	<p>It is the user's responsibility to set retention periods.</p> <p>TrackWise has an infinite retention period and data is retrievable at any time. If data is archived, the data can be reported upon in the archive system or sent back to the Production TrackWise system.</p>
<p>11.10(d)</p> <p>Is system access limited to authorized individuals?</p>	<p>7.1</p> <p>How is data secured by both physical and electronic means against damage? How is data accessible throughout the retention period?</p> <p>12.2</p> <p>The extent of security controls depends on the criticality of the system.</p>	<p>User & Sparta</p>	<p>The using organization is responsible for defining authorized access to the system.</p> <p>TrackWise allows for multiple configurable security groups, limiting access by process and by field.</p>
<p>11.10(e)</p> <p>Is there a secure, computer generated, time stamped audit trail that records the date and time of entries and actions that create, modify, or delete electronic records?</p>	<p>9</p> <p>Is an audit trail available to document the creation, change or deletion of data?</p> <p>12.4</p> <p>Is the system designed to record the identity of operators entering, changing, confirming or deleting data including date and time?</p>	<p>Sparta</p>	<p>TrackWise provides full audit trail for create and modify operations. Deletion of record data is not possible.</p> <p>The audit trail records the identity of operators entering, changing, confirming, or deleting data, including date and time.</p>

21 CFR Part 11	Annex 11	Responsible Party	TrackWise
11.10(e) Upon making a change to an electronic record, is the previously recorded information still available (e.g. not obscured by the change)?		Sparta	The TrackWise Audit Trail records previous values. Audit Trail entries cannot be deleted.
11.10(e) Is an electronic record's audit trail retrievable throughout the record's retention period?		Sparta	The audit trail is available for the life of the record.
11.10(e) Is the audit trail available for review and copying by the FDA?		Sparta	The TrackWise Activity History details are available for querying and reporting.
11.10(f) If the sequence of system steps or events is important, is this enforced by the system?		User & Sparta	TrackWise allows for fully configurable workflow management, thus the user can define the sequence of steps and events and ensure the proper process must be followed.
11.10(g) Does the system ensure that only authorized individuals can use the system, electronically sign records, access the operation, or computer system input or output device, alter a record, or perform other operations?	12.1 Are physical and/or logical controls in place to restrict access to the system?	User & Sparta	TrackWise allows for fully configurable security groups. The using organization needs procedures to define how application authorization is carried out. The using organization is responsible for restricting physical access.
11.10(h) If it is a requirement of the system that input data or instructions can only come from certain input devices (e.g. terminals) does the system check the validity of the source of any data or instructions received	4.8 When data is transferred to another data format or system, does the system check the validity to confirm data was not altered in value and/or meaning during migration.	Sparta	TrackWise checks that inputs are received in the browser in which the system was validated. Data migration tools ensure that no data was altered in value or meaning during migration from one TrackWise system to another TrackWise system

21 CFR Part 11	Annex 11	Responsible Party	TrackWise
<p>11.10(i)</p> <p>Is there documented training, including on the job training for system users, developers, IT support staff?</p>	<p>2</p> <p>Is there close cooperation between all relevant personnel such as process owner, system owner, qualified persons and IT?</p> <p>Do all personnel have appropriate qualifications, level of access and defined responsibilities to carry out their assigned duties?</p>	<p>User & Sparta</p>	<p>Within Sparta, employees are formally trained on policies, SOPs and work instructions. Employees also receive on the job training appropriate to their responsibilities. These SOPs outline how relevant personnel work together to complete their tasks and areas of responsibility.</p> <p>It is the using organization's responsibility to demonstrate that their staff has the education, training and experience to perform their assigned tasks.</p>
<p>11.10(j)</p> <p>Is there a written policy that makes individuals fully responsible for actions initiated under their electronic signatures?</p>		<p>User</p>	<p>This is the responsibility of the using organization.</p>
<p>11.10(k)</p> <p>Is the distribution of, access to, and use of systems operation and maintenance documentation controlled?</p>		<p>User</p>	<p>This is the responsibility of the using organization.</p>
<p>11.10(k)</p> <p>Is there a formal change procedure for system documentation that maintains a time sequenced audit trail of changes?</p>		<p>User & Sparta</p>	<p>It is the responsibility of using organization to ensure adequate change control procedures for documentation.</p> <p>Sparta Systems, Inc. maintains an audit trail for all system documentation and changes.</p>

Additional Procedures & Controls for Open Systems

21 CFR Part 11	Annex 11	Responsible Party	TrackWise
11.30 Is data encrypted?	5. Data What built-in checks are in place to confirm the correct and secure entry and processing of data?		Not Applicable. Closed System
11.30 Are digital signatures used?			Not Applicable. Closed System

Signed Electronic Records

21 CFR Part 11	Annex 11	Responsible Party	TrackWise
11.50 Do signed electronic records contain the following related information? <ul style="list-style-type: none"> • The printed name of the signer • The date and time of signing • The meaning of the signing (such as approval, review, responsibility) 	14 (c) Do electronic signatures include the time and date applied?	Sparta	Yes.
11.50 Is the above information shown on displayed and printed copies of the electronic record?		User & Sparta	It is the responsibility of the using organization to develop and validate reports.
11.70 Are signatures linked to their respective electronic records to ensure that they cannot be cut, copied, or otherwise transferred by ordinary means for the purpose of	14(b) Are electronic signatures permanently linked to their respective record	Sparta	Yes

21 CFR Part 11	Annex 11	Responsible Party	TrackWise
falsification?			

Electronic Signatures – General

21 CFR Part 11	Annex 11	Responsible Party	TrackWise
11.100(a) Are electronic signatures unique to an individual?	Not covered	User & Sparta	It is the responsibility of the using organization to ensure uniqueness to individual users. TrackWise enforces uniqueness by use of Person Identification Number (PID) generated by the database. Two logins of the same name and domain cannot exist.
11.100(a) Are electronic signatures ever reused by, or reassigned to, anyone else?	Not covered	User & Sparta	It is the responsibility of the using organization to ensure electronic signatures are not reused and/or reassigned to another user. TrackWise enforces uniqueness by not allowing duplicate login accounts and by use of PIDs.
11.100(b) Is the identity of an individual verified before an electronic signature is allocated?	Not covered	User	It is the using organization's responsibility to verify the identity of individuals assigned to an electronic record. Login to the system must occur by a named user before e-signature.
11.100(c) Can the user certify that the electronic signatures in their system are the legally binding equivalent to traditional handwritten signatures?	14 (a) Do electronic signatures have the same impact as hand-written signatures within the boundaries of the company?	User	It is entirely the responsibility of the customer to manage this certification to the agency.

Electronic Signatures – Non-Biometric

21 CFR Part 11	Annex 11	Responsible Party	TrackWise
11.200(a) (1) Is the signature made up of at least two components, such as an identification code and password?		Sparta	Signatures in TrackWise consist of a User ID and Password.
11.200(a) (1) (i) When several signings are made during a continuous session, is the password executed at each signing? Note: both components must be executed at the first signing of the session.		Sparta	The User ID and password are entered at each signing.
11.200(a) (1) (ii) If signings are not done in a continuous session, are both components of the electronic signature executed with each signing?		Sparta	The User ID and Password are entered at each signing.
11.200(a) (2) Are non-biometric signatures only used by their genuine owners?		User	It is the responsibility of the using organization to ensure employees only use their own electronic signature.
11.200(a) (3) Would an attempt to falsify an electronic signature require the collaboration of at least two individuals?		User	Using organizations need procedures that users do not divulge their electronic signature (e.g. password).

Electronic Signatures – Biometric

21 CFR Part 11	Annex 11	Responsible Party	TrackWise
11.200(b) Has it been shown that biometric electronic signatures can be used only by their genuine owner?			Not Applicable.

Controls for Identification Codes and Passwords

21 CFR Part 11	Annex 11	Responsible Party	TrackWise
11.300(a) Are controls in place to maintain the uniqueness of each combined identification code and password, such that no individual can have the same combination of identification code and password?		User & Sparta	It is the responsibility of the using organization to ensure uniqueness to individual users. TrackWise enforces uniqueness by use of Person Identification Number (PID) generated by the database.
11.300(b) Are procedures in place to ensure that the validity of identification codes is periodically checked?	11 Alterations to a system or to a computer program should only be made in accordance with a defined procedure which should include provision for validating, checking, approving and implementing the change. Such an alteration should only be implemented with the agreement of the person responsible for the part of the system concerned, and the alteration should be recorded. Every significant modification should be validated.	User	The management of change for a fully validated and deployed system is the sole responsibility of the customer. Sparta Systems may be contracted to assist in the deployment and validation of an approved change, but the customer is responsible for maintaining the Change Control process.

21 CFR Part 11	Annex 11	Responsible Party	TrackWise
11.300(b) Do passwords periodically expire and need to be revised?		User & Sparta	It is the responsibility of the using organization to set rules for expiration dates. TrackWise can be configured to enforce those procedures via "Days for Password Expiration" feature.
11.300(b) Is there a procedure for recalling identification codes and passwords if a person leaves or is transferred?	12.3 Is the creation, change and cancellation of access authorisations recorded?	User & Sparta	It is the responsibility of the using organization to establish procedures for recalling identification codes and passwords. TrackWise allows a user to be rendered inactive, without losing that user's historical activity. The modification of user access is recorded.
11.300(c) Is there a procedure for electronically disabling an identification code or password if it is potentially compromised or lost?	12.3 Is the creation, change and cancellation of access authorisations recorded?	User & Sparta	It is the responsibility of the using organization to establish procedures for disabling an identification code and/or password. TrackWise account can be set to inactive and have its password reset.
11.300(d) Is there a procedure for detecting attempts at unauthorized use and for informing security?		User & Sparta	It is the responsibility of the using organization to describe how to respond to attempted or actual unauthorized access. TrackWise will lock out user and provide notification after a specified number of failed attempts to login (set by configuration), execute an electronic signature or change a password.
11.300(d) Is there a procedure for reporting repeated attempts at unauthorized use of the system to management?		User & Sparta	It is the responsibility of the customer to provide a procedure for reporting repeated or serious attempts at unauthorized use. TrackWise can be configured to notify the administrator when a set number of login attempts in a single instance were unsuccessful.

Controls for Identification Codes and Passwords – For tokens, cards, and other devices bearing or generating identification code or password information

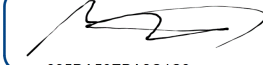
21 CFR Part 11	Annex 11	Responsible Party	TrackWise
11.300(c) Is there a loss management procedure to be followed if a device is lost or stolen?			Not applicable.
11.300(c) Is there a procedure for electronically disabling a device if it is lost, stolen, or potentially compromised?			Not applicable.
11.300(c) Are there controls over the issuance of temporary and permanent replacements?			Not applicable.
11.300(e) Is there an initial and periodic testing of tokens and cards?	11 Periodic Evaluation		Not applicable.
11.300(e) Does this testing check that there have been no unauthorized alterations?			Not applicable.

EU Annex 11 Controls for which there is no 21 CFR Part 11 Equivalent

Annex 11	Responsible Party	TrackWise
<p>1 Risk Management</p> <p>Are decisions on the extent of validation and data integrity controls based on a justified and documented risk assessment?</p>	User & Sparta	Using organizations are responsible for decisions regarding validation and data integrity controls.
<p>3.1</p> <p>When third parties are used to provide, install, configure, integrate, validate, maintain, modify or retain the system, do formal agreements exist?</p>	User & Sparta	<p>Using organizations are responsible for developing and executing agreements with third parties.</p> <p>Sparta maintains formal contracts with all third parties utilized for staff augmentation purposes.</p>
<p>3.1</p> <p>Do agreements with third parties clearly define the responsibilities of the third party?</p>	User & Sparta	<p>Using organizations are responsible for developing and executing agreements with third parties.</p> <p>Sparta maintains formal contracts with all third parties utilized for staff augmentation purposes.</p>
<p>3.2</p> <p>Are third parties audited?</p>	User & Sparta	<p>Using organizations are responsible for auditing any third parties they utilize.</p> <p>Sparta periodically audits all critical vendors.</p>
<p>3.3</p> <p>Is documentation from commercial off-the-shelf products reviewed to check that user requirements are fulfilled?</p>		Not applicable.
<p>3.4</p> <p>Is quality system and audit information relating to third party suppliers or developers of software & implemented systems available to inspectors on request?</p>		Not applicable.
<p>4.3</p> <p>Is an up to date listing of relevant systems and their GMP functionality available? For critical systems, an up to date system description detailing the physical and logical arrangements, data flows and interfaces with other systems or processes, hardware and software pre-requisites and security measures is available.</p>	User & Sparta	Using organizations are responsible for maintaining system lists and descriptions.

Annex 11	Responsible Party	TrackWise
<p>4.4</p> <p>Do user requirement specifications describe the required functions of the system? Is URS based on documented risk assessment and GMP impact. Are User requirements traceable throughout the life-cycle?</p>	Sparta	User requirement specifications drive system design and a traceability matrix is provided. User requirements are the responsibility of the using organization.
<p>4.5</p> <p>Was the system developed in accordance with an appropriate quality management system?</p>	Sparta	Sparta Systems is ISO 9001:2008 certified.
<p>4.6</p> <p>For customized systems, what process is in place to ensure the formal assessment and reporting of quality and performance measures for the life-cycle stages of the system.</p>		Not applicable. TrackWise is not customized.
<p>4.7</p> <p>What evidence of test methods and scenarios are available?</p> <p>Were parameter limits, data limits and error handling considered?</p> <p>How are automated testing tools and test environments assessed for adequacy?</p>	Sparta	<p>A validation package is available for each release.</p> <p>Parameter limits, data limits and error handling are considered during validation.</p> <p>Testing tools and environments use industry-leading tools whenever possible, and are otherwise reviewed for adequacy.</p>
<p>6 Accuracy Checks</p> <p>What accuracy checks are in place for critical data entered manually?</p>	User	Critical data fields can be configured to require the use of a drop-down selection list.
<p>7.2</p> <p>Are regular back-ups of relevant data done? How is the integrity and accuracy of data and the ability to restore data checked during validation and monitored periodically?</p>	User	Data back-ups are the responsibility of the using organization.
<p>8.2</p> <p>For records supporting batch release, are printouts available to indicate if any data was changed since original entry?</p>		Not applicable.
<p>10</p> <p>Are system changes made in a controlled manner in accordance with a defined procedure?</p>	User	Using organizations are responsible for defining a procedure for system changes. TrackWise maintains an audit trail of system changes

Annex 11	Responsible Party	TrackWise
<p>13</p> <p>Are all incidents reported and assessed? Is the root cause of critical incidents identified? Does the identified root cause form the basis of corrective and preventive actions?</p>	<p>User and Sparta</p>	<p>All product related incidents are brought to a weekly meeting where they are prioritized, severity noted and effort is decided. All high severity incidents are investigated, root cause analysis completed, and if applicable, a corrective action is identified.</p>
<p>15</p> <p>Does the system allow only qualified persons to certify the release of batches and clearly identify and record the person releasing or certifying the batches?</p>		<p>Not Applicable.</p>
<p>16</p> <p>What provisions are made to ensure continuity of support for critical processes in the event of a system breakdown?</p> <p>Is the time required to bring alternative arrangements into use based on risk and appropriate for the system and business process it supports?</p> <p>Are these arrangements adequately documented and tested?</p>		<p>The using organization is responsible for system uptime and redundancy and availability of backups</p>

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