

Internal Audit Check list

INTERNAL PROCESS VERIFICATION

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Company / ISO Section	Criteria of ISO Section	Auditor Comments / Issues
VST Ltd ISO9001:201 5.5.1.1	<p>General</p> <p>Top management shall demonstrate leadership and commitment with respect to the quality management system by:</p> <ul style="list-style-type: none"> a) taking accountability for the effectiveness of the quality management system; b) ensuring that the quality policy and quality objectives are established for the quality management system and are compatible with the context and strategic direction of the organization; c) ensuring the integration of the quality management system requirements into the organization's business processes; d) promoting the use of the process approach and risk-based thinking; e) ensuring that the resources needed for the quality management system are available; f) communicating the importance of effective quality management and of conforming to the quality management system requirements; g) ensuring that the quality management system achieves its intended results; h) engaging, directing and supporting persons to contribute to the effectiveness of the quality management system; i) promoting improvement; j) supporting other relevant management roles to demonstrate their leadership as it applies to their areas of responsibility. <p>NOTE Reference to "business" in this International Standard can be interpreted broadly to mean those activities that are core to the purposes of the organization's existence, whether the organization is public, private, for profit or not for profit.</p>	/
VST Ltd ISO9001:201 5.5.2.1	<p>Establishing the quality policy</p> <p>Top management shall establish, implement and maintain a quality policy that:</p> <ul style="list-style-type: none"> a) is appropriate to the purpose and context of the organization and supports its strategic direction; b) provides a framework for setting quality objectives; c) includes a commitment to satisfy applicable requirements; d) includes a commitment to continual improvement of the quality management system. 	/

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VST Ltd ISO9001:201 5.6.2.2	When planning how to achieve its quality objectives, the organization shall determine: a) what will be done; b) what resources will be required; c) who will be responsible; d) when it will be completed; e) how the results will be evaluated.	
VST Ltd ISO9001:201 5.7.5.1	General 7.5.1 General The organization's quality management system shall include: a) documented information required by this International Standard; b) documented information determined by the organization as being necessary for the effectiveness of the quality management system. NOTE The extent of documented information for a quality management system can differ from one organization to another due to: — the size of organization and its type of activities, processes, products and services; — the complexity of processes and their interactions; — the competence of persons.	
Viamed Ltd ISO13485:20 16.4.1.3	Quality management system For each quality management system process, the organization shall: a) determine criteria and methods needed to ensure that both the operation and control of these processes are effective; b) ensure the availability of resources and information necessary to support the operation and monitoring of these processes; c) implement actions necessary to achieve planned results and maintain the effectiveness of these processes; d) monitor, measure as appropriate, and analyse these processes; e) establish and maintain records needed to demonstrate conformance to this International Standard and compliance with applicable regulatory requirements (see 4.2.5).	Roles + titles management Review Rolling issues Doc index
Viamed Ltd ISO13485:20 16.4.1.4	Quality management system For each quality management system process, the organization shall: The organization shall manage these quality management system processes in accordance with the requirements of this International Standard and applicable regulatory requirements. Changes to be made to these processes shall be: a) evaluated for their impact on the quality management system; b) evaluated for their impact on the medical devices produced under this	Roles + titles management Review Doc index

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	quality management system c) controlled in accordance with the requirements of this International Standard and applicable regulatory requirements.	
Viamed Ltd ISO13485:20 16.4.2.1 General	Documentation requirements The quality management system documentation (see 4.2.4) shall include: a) documented statements of a quality policy and quality objectives; b) a quality manual; c) documented procedures and records required by this International Standard; d) documents, including records, determined by the organization to be necessary to ensure the effective planning, operation, and control of its processes; e) other documentation specified by applicable regulatory requirements.	Doc index Route map management Review
Viamed Ltd ISO13485:20 16.4.2.2 Quality manual	Documentation requirements The organization shall document a quality manual that includes: a) the scope of the quality management system, including details of and justification for any exclusion or non-application; b) the documented procedures for the quality management system, or reference to them; c) a description of the interaction between the processes of the quality management system. The quality manual shall outline the structure of the documentation used in the quality management system.	Doc index Route map Procedure
Viamed Ltd ISO13485:20 16.5.1	Management commitment Top management shall provide evidence of its commitment to the development and implementation of the quality management system and maintenance of its effectiveness by: a) communicating to the organization the importance of meeting customer as well as applicable regulatory requirements; b) establishing the quality policy; c) ensuring that quality objectives are established; d) conducting management reviews; e) ensuring the availability of resources.	Procedures meetings Roles and titles management Review Doc index
Viamed Ltd ISO13485:20 16.5.4.1	Quality objectives Top management shall ensure that quality objectives, including those needed to meet applicable regulatory requirements and requirements for product, are established at relevant functions and levels within the organization. The quality objectives shall be measurable and consistent with the quality policy.	management Review Route map Doc index Procedures Roles + tasks

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Viamed Ltd ISO13485:20 16.5.4.2	Quality management system planning Top management shall ensure that: a) the planning of the quality management system is carried out in order to meet the requirements given in 4.1, as well as the quality objectives; b) the integrity of the quality management system is maintained when changes to the quality management system are planned and implemented.	Management Review Route maps procedures Roles + titles
Viamed Ltd ISO13485:20 16.5.5.1	Responsibility and authority Top management shall ensure that responsibilities and authorities are defined, documented and communicated within the organization. Top management shall document the interrelation of all personnel who manage, perform and verify work affecting quality and shall ensure the independence and authority necessary to perform these tasks.	Roles and titles Issues Doc index management Review
Viamed Ltd ISO13485:20 16.5.5.2	Management representative Top management shall appoint a member of management who, irrespective of other responsibilities, has responsibility and authority that includes: a) ensuring that processes needed for the quality management system are documented; b) reporting to top management on the effectiveness of the quality management system and any need for improvement; c) ensuring the promotion of awareness of applicable regulatory requirements and quality management system requirements throughout the organization.	Roles + titles Issues Doc index management Review Procedures
Viamed Ltd ISO13485:20 16.5.6.3	Review output The output from management review shall be recorded (see 4.2.5) and include the input reviewed and any decisions and actions related to: a) improvement needed to maintain the suitability, adequacy, and effectiveness of the quality management system and its processes; b) improvement of product related to customer requirements; c) changes needed to respond to applicable new or revised regulatory requirements; d) resource needs.	Doc index QA system management Review Issues
Viamed Ltd ISO13485:20 16.6.1	Provision of resources The organization shall determine and provide the resources needed to: a) implement the quality management system and to maintain its effectiveness; b) meet applicable regulatory and customer requirements.	Route maps Issues Procedures
Viamed Ltd ISO13485:20	Internal audit The organization shall conduct internal audits at planned intervals to	

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16.8.2.4	<p>determine whether the quality management system:</p> <p>a) conforms to planned and documented arrangements, requirements of this International Standard, quality management system requirements established by the organization, and applicable regulatory requirements;</p> <p>b) is effectively implemented and maintained.</p> <p>The organization shall document a procedure to describe the responsibilities and requirements for planning and conducting audits and recording and reporting audit results.</p> <p>An audit program shall be planned, taking into consideration the status and importance of the processes and area to be audited, as well as the results of previous audits. The audit criteria, scope, interval and methods shall be defined and recorded (see 4.2.5). The selection of auditors and conduct of audits shall ensure objectivity and impartiality of the audit process.</p> <p>Auditors shall not audit their own work.</p> <p>Records of the audits and their results, including identification of the processes and areas audited and the conclusions, shall be maintained (see 4.2.5).</p> <p>The management responsible for the area being audited shall ensure that any necessary corrections and corrective actions are taken without undue delay to eliminate detected nonconformities and their causes. Follow-up activities shall include the verification of the actions taken and the reporting of verification results.</p> <p>NOTE Further information can be found in ISO 19011.</p>	<i>Doc index</i> <i>Audit calendar</i> <i>Role and titles</i> <i>Ranking</i>
Viamed Ltd ISO13485:20 16.8.3.4	<p>Rework</p> <p>The organization shall perform rework in accordance with documented procedures that takes into account the potential adverse effect of the rework on the product. These procedures shall undergo the same review and approval as the original procedure.</p> <p>After the completion of rework, product shall be verified to ensure that it meets applicable acceptance criteria and regulatory requirements.</p> <p>Records of rework shall be maintained (see 4.2.5).</p>	<i>Procedures</i> <i>QA systems</i> <i>Doc index</i>
Viamed Ltd ISO13485:20 16.8.5.3	<p>Preventive action</p> <p>The organization shall determine action to eliminate the causes of potential nonconformities in order to prevent their occurrence. Preventive actions shall be proportionate to the effects of the potential problems.</p> <p>The organization shall document a procedure to describe requirements for:</p> <p>a) determining potential nonconformities and their causes;</p> <p>b) evaluating the need for action to prevent occurrence of nonconformities;</p>	<i>Doc index</i> <i>Procedures</i> <i>Management Review</i>

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	<p>c) planning and documenting action needed and implementing such action, including, as appropriate, updating documentation;</p> <p>d) verifying that the action does not adversely affect the ability to meet applicable regulatory requirements or the safety and performance of the medical device;</p> <p>e) reviewing the effectiveness of the preventive action taken, as appropriate.</p> <p>Records of the results of any investigations and of action taken shall be maintained (see 4.2.5).</p>	<i>Review meetings</i>
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	INTERNAL PROCESS VERIFICATION A. Management System: B. Management Responsibility C. Resource Management D. Product Realisation E. Design & Development F. Product Provision G. Process Monitoring The following are questions that should be asked and answered either through Internal audits or at this meeting		
1	Review Last years Audit. Update processes if required. Are all follow on Issue resolved satisfactory.	<i>Action outstanding</i>	Y
2	Is the Quality Statement Policy and Objectives reviewed annually. ISO – Document Index Task ID (300). Search Issues and review.	<i>262913 ✓</i>	Y
3	Is the process manual up to date. ISO – Document Index Task ID (548). Search Issues and review.	<i>275145 ✓</i>	Y
4	Is VOP / VOP documentation checked prior to formal approval and issue.		Y
5	Check that there is a system in operation for the request for amendments.	<i>See update request</i>	Y
6	Verify that amendments are updated electronically and old copies archived.		Y
7	Are sales orientated records filed and archived correctly.	Digital and Automatic attached to contact order	Y

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V16M201

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8	Has organisation Chart changed. VM3COP02.02		✓
9	Has personnel responsibility descriptions changed. Roles Titles Processes and Procedures ADMIN Over View for complete list		✓
10	Check that the CE files are maintained by sole responsibility.		✓
11	Check that the Notified body is informed of major changes to Documentation.	in changes	✓
12	Check that electronic documents are regularly backed up and secure off site. ISO – Document Index Task ID (452). Search Issues and review.	276728 ✓	✓
13	Does the management system comprise a series of process controls and are they in place throughout the organisation. Are processes identified.	Intrastats, Audit 10 Roles + titles	✓
14	Check the system for its policies and objectives and its control of the above processes and procedures. Is the Standards Manual up to date and does it indicates the company's objectives. Are procedures in place - VM3COP Are they available to all personnel – Doc Index Are other company documents i.e. Technical Drawings, Standards; Operators Manuals etc. also available and controlled Doc index	Intrastats, Audit 10 Roles and Responsibilities.	✓
15	Are the latest revision of documents controlled by version and date status and are they easily accessible. Is the Managing Director or designate manager still giving final approval for document changes.	Intrastats, Audit 10	✓
16	Has the Business Continuity Plan has expired. ISO – Document Index Task 266.	250% ✓	✓
<u>B - MANAGEMENT RESPONSIBILITY</u>			
17	Is Top management showing full commitment to the overall system and are communication lines in place. Manage Review Task 290.	Intrastats, Director in control of QA system 277403 ✓	✓
18	Are all customer requirements defined and met.	Contract Review Audit 2	✓
19	Are all the processes and objectives, undertaken within the		

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	company, documented in Intrastats and have a procedure. Is it measurable. Check process for measurable task 114 Documented in Staff – Audit of Roles, titles and procedures.	276867 ✓	✓
20	Does the person responsible for the management systems have the authority to implement actions and reports directly to top management with the need for these actions	Managing Director	✓
21	Are reviews of the management system undertaken regularly and the results and actions relayed throughout the organisation. Task 290 for weekly review 277403 ✓ Task 114 for bigger overview 276867 ✓ Task 746 for total review 267637 ✓	Issues, Message of Day, company meetings, management meetings, Management weekly reviews	✓
22	Are all required actions undertaken in a timely manner and closed where appropriate.	Intrastats Issues	✓
23	Are all output requirements in such a format that verification against inputs, is applicable and appropriate. Is fitness for Purpose validated and is it measurable. Staff – Audit of Roles, titles and procedures – click into details - review Scope and Risks. To check relevance. Staff – Audit of Roles, titles and procedures check down the page for gaps in the IP 1-6 (end tick boxes)		✓
24	Are actions recorded against verifications completed in a timely and responsible manner.	Intrastats Issues	✓
25	Are design changes recorded and all the relevant information filed in the appropriate places.	Design control Audit 3 Intrastats 2020-01-01	✓
C - RESOURCE MANAGEMENT			
26	Has top management established a mechanism for identifying and providing required resources, training etc.	Training Audit 8	✓
27	Does this include existing and new personnel.	Training Audit 8	✓
28	Has top management identified the competency levels and attributes required for existing and new personnel.	Training Audit 8	✓
29	Is the competency of personnel monitored, verified and the appropriate records maintained	Training Audit 8	✓
30	Are personnel responsibilities defined.	Roles and Responsibilities	✓

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31	Do individuals know their responsibilities, reporting and communicating lines. Each employee has 'My Roles' Link Task 314	Intrastats communication 270335 ✓	✓
32	Verify that all procedures, detail who is responsible for it.		✓
33	Check these roles and responsibilities also include Health & Safety tasks – Health and Safety Controller.		✓
34	Is the need for equipment, plant, services etc. identified and acted upon where necessary. Task 13 Part of audit 19	267572 ✓ Production meetings, management meetings Health and Safety Questionnaire.	✓
35	Has the basic working infrastructure been planned with conformity to requirements in mind.	Health & safety Audit 19	✓
36	Are there adequate mechanisms in place for the identification, handling etc. of product through all stages.	COP/07	✓
37	Are the controls in place, to safeguard customer property, adequate for full protection against loss damage etc.	COP/09	✓
38	Is the process for monitoring and measurement of product in place at all stages throughout the production process.	Production COPs	✓
39	Is the process for control of measuring equipment adequate for the monitoring of product verifications.	Calibration Audit 06	✓
40	Are validity processes in place to safeguard product integrity.	Bar coding traceability	✓
D - PRODUCT REALISATION			
41	Is the planning process for the realisation of product undertaken at the relevant stages.		✓
42	Does planning identify documentation, testing and other such activities as required and that all appropriate records are maintained.		✓
43	Are all customer requirements being addressed, including statutory and regulatory and that the capabilities are identified to meet those requirements.	Contract Review Audit 02	✓
44	Establish that mechanisms are in place to review all customer requirements prior to any commitments by the organisation.	Contract Review Audit 02	✓
45	Check that there are adequate arrangements for customer communications and feedback.	Contract Review Audit 02	✓
46	Is collation and analysis of all relevant data determined and effective. Is corrective actions identified.		✓
47	Are these actions completed in a timely and adequate manner and are these actions part of continual improvements.		✓
48	Does the organisation have preventive measures in place to		✓

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	control potential non-conformities.		
49	Are all the above actions are reviewed adequately.	No Nee Design	Y
E - DESIGN & DEVELOPMENT			
50	Are procedures in place to ensure adequate planning of product design and that all relevant stages are identified.	Design control Audit 3	N/A
51	Are the interfaces and assignments of responsibilities identified.	Design control Audit 3	N/A
52	Are all input requirements determined. Is the documentation identified.	Design control Audit 3	N/A
53	Are all output requirements in such a format, that verification against inputs is applicable and appropriate. Is Fitness for Purpose validated.	Design control Audit 3	N/A
54	Are actions recorded against verifications completed in a timely and responsible manner.	Design control Audit 3	N/A
55	Are validation processes in place and are they determined in accordance with the relevant requirements.	Design control Audit 3	N/A
56	Are design changes recorded and all the relevant information filed in the appropriate places.	Design control Audit 3	N/A
F - PRODUCT PROVISION			
57	Are supplier profiles adequate and appropriate for the organisation. Are they monitored, for their ability to provide the requirements, is this monitoring controlled.	Purchasing Controls (Supplier Performance) Audit 5	Y
58	Is all the required information necessary, forwarded to suppliers in the correct format. Will be system controlled.	Purchasing Controls (Supplier Performance) Audit 5	Y
59	Are goods and services received correct to the requirements stipulated.	Goods Inward Audit 9	Y
60	Are the provisions available, suitable for control of production and service, including procedures and equipment etc.	Production Audit 15	Y
61	Are there adequate mechanisms in place for the identification, handling etc. of product through all stages.	Production Audit 15	Y
62	Are the controls in place to safeguard customer property adequate for full protection against loss damage etc.	Production Audit 15	Y
63	Is the process for monitoring and measurement of products in place at all stages throughout the production process.	Production Audit 15	Y
64	Is the process, for control of measuring equipment, adequate for the monitoring of product verifications.	Calibration Audit 6	Y
G - PROCESS MONITORING			
65	Are mechanisms in place to monitor all relevant processes,		

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	including customer satisfaction. Are these verified against known criteria. Check process ID 114	276867 ✓	✓
66	Are controls in place for non-conforming product and processes. Are adequate to prevent unintended uses.	Goods Inward Audit 9	✓
67	Where non-conforming product / process have been detected is appropriate action taken.	Goods Inward Audit 9	✓
68	Is collation and analysis of all relevant data determined and effective Is corrective actions identified.		✓
69	Are these actions completed in a timely and adequate manner. Are these actions part of continual improvements.		✓
70	Are all the above actions are reviewed adequately. Check process ID 114	Annually 276867 ✓	✓
71	Are regular analyses undertaken to identify any outstanding requirements.	Intrastats	✓
72	Are necessary changes implemented where and when required.		✓
73	Is any outsourcing done.		No
74	Check the documented system for its policies, objectives and its control of the above processes and procedures. Intrastats – document index – VM3COP00.00 / VM3COP00.01. Check documents for location of objectives and policies.	Intrastats	✓
75	Are records of inspections filed.	Audits	✓

List Processes Per Title

Managing Director

Process Scope	Roll Task	Roll Audit	Risk	Action	Referenced in Document
PROCESSID 7837 To Review the External Parties Influencing The QMS VST / Viamed Checked the Scopes and Risks. Review the Underlining Processes and Tasks	743 Managing Director	784 Company Secretary	Freq 1 Risk 1 Overall 1	Task 12M Audit 12M	
PROCESSID 7845 Determine, provide and maintain the environment necessary for the operation of its processes and to achieve conformity of products and services.	745 273>01	276334	Freq 1 Risk 1 Overall 1		

Merged into 7729 can close the tasks

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PROCESSID 7846

To Comply with Top Level
Re-authorise the Current Audits for next 12 Months
Cover the Agenda as Per VOP13

746 Freq 1 Task 12M

Managing Director Risk 1 Overall 1

267657+ ongoing

PROCESSID 7848

To Review the Scope of the
ISO 9001 / ISO 13485 Standards

749 Freq 1 Task 12M

Managing Director Risk 1 Overall 1

272935 ✓

PROCESSID 7871

To review the Exclusions / boundaries to ISO
13485:2016 for Viamed

790 Freq 1 Task 12M

Managing Director Risk 1 Overall 1

267641

ISO Controller
Process Scope

Roll Task **Roll Audit** **Risk** **Action** **Referenced in Document**

PROCESSID 6866

Review the Internal Process and Verification's are suitable for the current standards

55 Freq 1 Risk 1 Overall 1

PROCESS NOW CANCELLED AS REPEAT OF AUDIT 20
PROCESSID 7827

To review the Quality policy and check it is still valid and upto date.

301 Freq 1 Task 12M

Managing Director Risk 1 Overall 1

269914 ✓

PROCESSID 7828

To review the Quality policy and check it is still valid and upto date.

723 Freq 1 Task 12M

Managing Director Risk 1 Overall 1

269947 ✓

IT Controller
Process Scope

Roll Task **Roll Audit** **Risk** **Action** **Referenced in Document**

PROCESSID 7701

Amazon Web Services, is an online service, which basically simply provides a Linux PC out on the Web. Viamed uses this, for Web development of Websites. It hosts a working backup of many websites. Viamed / vst / vandagraph etc..

511 Freq 3 Task 1M

Office Processes Risk 1 Overall 3

276890 ✓ terms

PROCESSID 7755

To Send Invoice for online services to Helen

597 Freq 3 Task 1M

Office Processes Risk 1 Overall 3

277060 ✓ terms

PROCESSID 7832

Backup of all Sent Emails sent to External Address for Verification

731 Freq 1 Task 2W

Managing Director Risk 1 Overall 1

277318 ✓

PROCESSID 7850

Test the Goods out process disabling picking of items not Goods Out

752 Freq 1 Task 9M

Managing Risk 1 Audit 12M

270649 249739 ✓

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relating to an order

PROCESSID 7851

To test intrastats does not allow picking of unprocessed products to live customer orders

754 Goods Out

755 Managing Director
262817

Overall 1

Freq 1 Task 6M
Risk 1 Audit 12M
Overall 1

PROCESSID 7852

To attempt to Scan a product that has gone past its expire date.

756 Goods Out

757 Managing Director
27436517

Freq 1 Task 12M
Risk 1 Audit 12M
Overall 1

PROCESSID 7853

Warehouse shelves can be tagged as sellable stock / unsellable stock. Either for quarantine purposes or holding items for other customer orders.

759 Goods Out

760 Managing Director
270652

Freq 1 Task 12M
Risk 3 Audit 12M
Overall 3

244103

Test that Order picking cannot pick unsellable stock locations to an Order

PROCESSID 7854

Software Validation of the production lists.

761 Goods In

762 Managing Director
273503

Freq 2 Task 3M
Risk 2 Audit 6M
Overall 4

276343

By confirming no extra production jobs are stuck in the system, and all listed production jobs are found, the production tracking is validated

761 Goods In

762 Managing Director
273503

Freq 2 Task 3M
Risk 2 Audit 6M

Overall 4

276343

PROCESSID 7855

Software Validation - Production Lists

Review the current active production lists in intrastats to the actual in progress production lists

761 Processes

762 Managing Director
273503

Freq 2 Task 3M
Risk 2 Audit 6M

Overall 4

276343

PROCESSID 7856

To check order picking cannot pick against an unchecked order

764 Office

765 Managing Director
246827

Freq 2 Task 12M
Risk 2 Audit 12M

Overall 4

276343

Processes

766 Director

Freq 2 Task 12M
Risk 2 Audit 12M

Overall 2

276343

PROCESSID 7857

To confirm Software Validation Stock Tracking Check, is functioning as expected

763 Goods In

1155 Managing Director
27410817

Freq 2 Task 6M
Risk 1 Audit 12M

Overall 2

276343

PROCESSID 7858

Test the QA System that Staff not trained for QA are unable to QA a Product.

766 Office

1175 Managing Director
261868

Freq 1 Task 6M
Risk 3 Audit 12M

Overall 3

276343

Processes

767 Director

Freq 1 Task 6M
Risk 2 Audit 12M

Overall 2

276343

PROCESSID 7861

Software Validating Of Training Documents via Forced Required Reading

768 Managing Director

Freq 1 Task 12M
Risk 2 Audit 12M

Overall 2

276343

PROCESSID 7865

Software Validation of the system:

To check all process(s) tasks and audits are not clashed with the same person doing the Task as the Audit.

779 Managing Director

781 Managing Director
27410

Freq 1 Task 12M
Risk 1 Audit 12M

Overall 1

276343

Processes

780 Director

Freq 1 Task 12M
Risk 1 Audit 12M

Overall 1

276343

Processes

PROCESSID 7870

Scope to check the automatic system of tagging product non conformance and other issues to the post market surveillance review report.

782 Managing Director

Freq 1 Task 12M
Risk 1 Audit 12M

Overall 1

276343

Processes

PROCESSID 7875

802

803

Freq 1 Task 12M

Overall 1

276343

Processes

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To test document control is working as intended.	Managing Director	Company Secretary	Risk 1	Audit 12M
PROCESSID 7879	808	809	Freq 1	Task 12M
To check the Scheduled Tasks and Audits is working as Intended.	Managing Director	Company Secretary	Risk 1	Audit 6M
To also Check the Out of Date documents is working as Intended.	248861	275592	Overall 1	
PROCESSID 7880	808	809	Freq 1	Task 12M
To confirm the out of documents computer software functions as expected flagging out of date items on to the list	Managing Director	Company Secretary	Risk 1	Audit 6M
PROCESSID 7881	810		Freq 1	
To compare Opera Live Orders to Intrastats Back order Active List			Risk 1	
			Overall 1	

NO LONGER REQUIRED

Opera is now out of the system

Audits

Process Scope	Roll Task	Roll Audit	Risk	Action	Referenced in Document
PROCESSID 7723		3			
To carry out Audit 10b Process Verification Viamed			Freq 1		
Now Defunct - See Audit 20			Risk 2		
PROCESSID 7730			Overall 2		
To carry out Audit 20 Process Verification To Management Viamed	276292	172	Freq 1	Audit 12M	
	This Audit	Company Secretary	Risk 2		
PROCESSID 7771		177	Overall 2		
To carry out Audit 10b Process Verification VST			Freq 1		
Now Defunct - See Audit 20			Risk 2		
PROCESSID 7778	276297	181	Freq 1	Audit 12M	
To carry out Audit 20 Process Verification To Management VST	VST Audit	Company Secretary	Risk 2		
			Overall 2		