

Internal Audit Check List

SERVICE LOGS	
Created:	17/May 1995
Revised:	12 June 2017
Audit Date	28-8-19 Auditor Helen Lamb

Company / ISO Section	Criteria of ISO Section	Auditor Comments / Issues
VST Ltd ISO9001:2015 8.5.1	<p>Control of production and service provision</p> <p>The organization shall implement production and service provision under controlled conditions. Controlled conditions shall include, as applicable:</p> <p>a) the availability of documented information that defines:</p> <ol style="list-style-type: none"> the characteristics of the products to be produced, the services to be provided, or the activities to be performed; the results to be achieved; the availability and use of suitable monitoring and measuring resources; the implementation of monitoring and measurement activities at appropriate stages to verify that criteria for control of processes or outputs, and acceptance criteria for products and services, have been met; the use of suitable infrastructure and environment for the operation of processes; the appointment of competent persons, including any required qualification; the validation, and periodic revalidation, of the ability to achieve planned results of the processes for production and service provision, where the resulting output cannot be verified by subsequent monitoring or measurement; the implementation of actions to prevent human error; the implementation of release, delivery and post-delivery activities 	<p>Doc notes</p> <p>In progress</p>

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Viamed Ltd ISO13485:2011 6.7.5.3	<p>Installation activities</p> <p>The organization shall document requirements for medical device installation and acceptance criteria for verification of installation, as appropriate.</p> <p>If the agreed customer requirements allow installation of the medical device to be performed by an external party other than the organization or its supplier, the organization shall provide documented requirements for medical device installation and verification of installation.</p> <p>Records of medical device installation and verification of installation performed by the</p>	<i>Calibration Audit Installations</i>
		<i>Acc. to clx No 3rd Party Installers/Service Providers</i>

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Viamed Ltd ISO13485:201 6.7.5.4	<p>organization or its supplier shall be maintained (see 4.2.5).</p> <p>Servicing activities</p> <p>If servicing of the medical device is a specified requirement, the organization shall document servicing procedures, reference materials, and reference measurements, as necessary, for performing servicing activities and verifying that product requirements are met.</p> <p>The organization shall analyse records of servicing activities carried out by the organization or its supplier:</p> <ul style="list-style-type: none"> a) to determine if the information is to be handled as a complaint; b) as appropriate, for input to the improvement process. <p>Records of servicing activities carried out by the organization or its supplier shall be maintained (see 4.2.5).</p> <p>Validation of processes for production and service provision</p> <p>The organization shall validate any processes for production and service provision where the resulting output cannot be or is not verified by subsequent monitoring or measurement and, as a consequence, deficiencies become apparent only after the product is in use or the service has been delivered.</p> <p>Validation shall demonstrate the ability of these processes to achieve planned results consistently.</p> <p>The organization shall document procedures for validation of processes including:</p> <ul style="list-style-type: none"> a) defined criteria for review and approval of the processes; b) equipment qualification and qualification of personnel; c) use of specific methods, procedures and acceptance criteria;
Viamed Ltd ISO13485:201 6.7.5.6	<p><i>Dec include</i></p> <p><i>management review</i></p> <p><i>instruments</i></p> <p><i>procedures</i></p>

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<p>d) as appropriate, statistical techniques with rationale for sample sizes</p> <p>e) requirements for records (see 4.2.5);</p> <p>f) revalidation, including criteria for revalidation;</p> <p>g) approval of changes to the processes.</p> <p>The organization shall document procedures for the validation of the application of computer software used in production and service provision. Such software applications shall be validated prior to initial use and, as appropriate, after changes to such software or its application. The specific approach and activities associated with software validation and revalidation shall be proportionate to the risk associated with the use of the software including the effect on the ability of the product to conform to specifications.</p> <p>Records of the results and conclusion of validation and necessary actions from the validation shall be maintained (see 4.2.4 and 4.2.5).</p>	<i>QA & Records</i> <i>Management Review</i>
<p>Viamed Ltd ISO13485:2011 6.8.2.4</p> <p>The organization shall conduct internal audits at planned intervals to 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</p>	<i>Analysts</i> <i>Planning Analysts</i> <i>Analysts</i>

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<p>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. 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>	<p>Analysis of data</p> <p>ISO13485:2011</p> <p>6.8.4</p> <p>The organization shall document procedures to determine, collect and analyse appropriate data to demonstrate the suitability, adequacy and effectiveness of the quality management system. The procedures shall include determination of appropriate methods, including statistical techniques and the extent of their use.</p> <p>The analysis of data shall include data generated as a result of monitoring and measurement and from other relevant sources and include, at a minimum, input from:</p> <ul style="list-style-type: none"> a) feedback; b) conformity to product requirements; c) characteristics and trends of processes and product including opportunities for improvement; d) suppliers; 	<p><i>Management Review + Internal Audit</i></p>
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e) audits;
f) service reports, as appropriate.
If the analysis of data shows that the quality management system is not suitable, adequate or effective, the organization shall use this analysis as input for improvement as required in 8.5.
Records of the results of analyses shall be maintained (see 4.2.5).

Internal Audit

Answer questions 1-4, Any non conformance generate an Issue.

INTRASTATS STOCK MENU

SERVICE VISITS

Service Visits

All Active and Single visit services should be shown.

1. Are Any service Visits Over Due?

Note Tracking ID 34 in the Example,

No Services Due

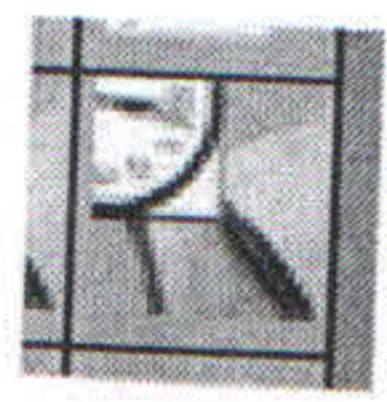
Tracking ID	Service ID	Recur Month	Opeta	Location	Description	Equipment List	Issue Date	Scheduled Date	ORDS	KEY	
										Recurring Service	Once Only Service
34	12	12	00004990	Sunderland Royal Hospital	Delivery Suite and Neonatal Unit	48 Items	26/07/15	...	ORD6051	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>
33	9	12	00004260	Royal Preston Hospital	Resus cabinets, Delivery Suite	41 Items	03/04/15	...		<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>
35	29	12	00000780	Burnley General Hospital	Resus Cabinets and Blenders	28 Items	11/09/15	...		<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>
36	27	12	00000550	Royal Blackburn Hospital	Resus cabinet and blender annual service	12 Items	17/09/15	...		<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>
14	14	0	00005210	Walsall Manor Hospital	Tron Thumb Conversions and Upgrade	4 Items	15/02/13	...		<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>
13	13	0	00001350	County Durham & Darlington Hospital	maternity tnm thumbs	10 Items	18/04/13	...		<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>
28	28	0	00000591	Royal Bolton Hospital	Resus cabinet upgrade to include blender	32 Items	17/05/13	...		<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>
25	18	0	00002370	Westmorland General Hospital	Resus cabinet upgrade to include blender	1 Items	19/06/14	...		<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>
30	23	0	00003580	North Manchester General Hosp	Resus cabinet and blender annual service	16 Items	30/07/14	...		<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>

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Has the Over due Visit got any Action notes:

Click:



No Overdue
Visits

2. Have any over due Visit NOT got any action notes?

Service Visit ID: 12 Tracking ID: 34

Location Notes

Hotel Notes

Dates Booked

Booked in Bede Guest House from Monday 26th January for three nights. Contact name at Guest House is Pamela Tate - 07833787481
Emailed Pam to advise that we Week commencing 26th January Booked in Bede Guest House from Monday 26th Jan for three nights. Had to cancel above dates as per Phil., will re-organize once Phil is feeling better.
Need to call/e-mail David Ferguson on the morning to OK the visit.

Engineer notes

Completion notes

Service Engineer has completed the visit
Service Engineer has scanned all relevant parts
Service Reports Sent
Primary Contact Sent Reports
Service Reports Confirmed Received
Invoice can now be generated/sent
Service Visit is fully complete

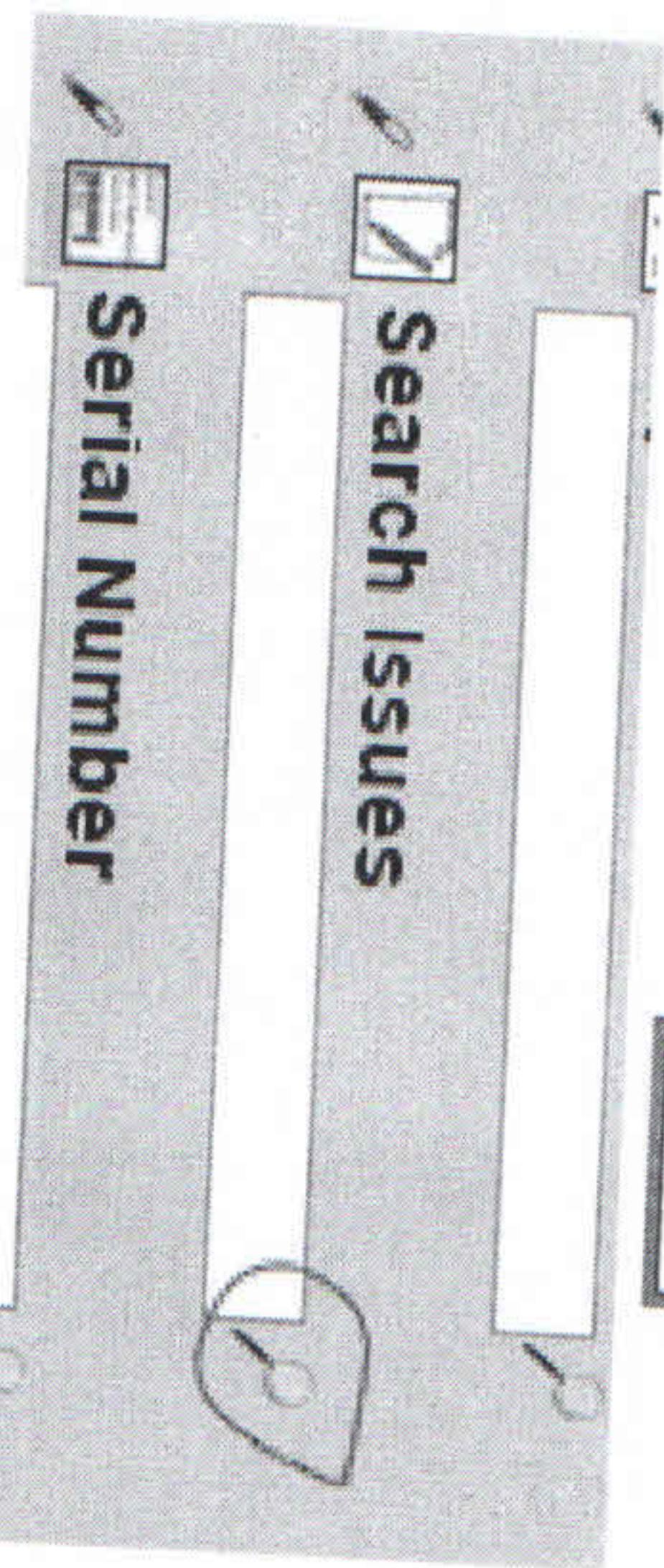
will flag ORD69051 Yellow in the Active List

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Drop out of the Service Visit Section,

Search Intrastats



In the Subject Contains Words box type: *Check The Service Visit Logs*

Done (not issue # 152248 And not # 152247 task ✓

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You should see a list of system generated tasks to 2 Different employees,
Scroll down the list,

3. Are more than 1 Issue outstanding per user?

No

4. Any problems found in the Issues system not being addressed?

No Issues *None*

Sub Processes Linked to Audit

Review the below processes tasks and audits and ensure they are completed in a timely manner.

32 Issues Found

Issue #	Description	Created By	Date Created	Completed By	Date Completed
53586	Service existing	Auto Calender	28/10/14		
53577	Service existing	Auto Calender	27/10/14	Lisa Leggoe	28 Oct 2014
53437	Service existing	Auto Calender	21/10/14		

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Audits

150658/ This Audit

Process Scope	Brief Description	Responsibility/Procedure/Training	Measurable Objective	Task	Audit	Freq	Risk	Overall Action
7889 - History/Details	Audit 24 To carry out Audit 24 Servicing Vianed	27178 VOP 13 Process Monitoring, System Reviews, Audits, Management Review, Analysis Data			288 Company Secretary	1	2	Audit 12M

Office Processes

Process Scope	Brief Description	Responsibility/Procedure/Training	Measurable Objective	Task	Audit	Freq	Risk	Overall Action
5857 - History/Details	Ensuring customer onsite service visits are completed	Customer Service Logs 24730 VOP 03 Contract Review, Enquires, Office Processes	233 Office Processes	234 UK Sales Controller	1	4	1	Task 1W Audit 1M
7760 - History/Details	Send letters to existing customers to remind them that a service is due on their equipment	Send Service Offers 24730 VOP 03 Contract Review, Enquires, Office Processes	607 Marketing Processes	898 Company Secretary	1	1	1	Task 1W Audit 4W