

Internal Audit Check list			
VIAMED LTD SERVICE LOGS			
Created:	17/May 1995	Audit No 24	
Audit Date	16-8-25	Auditor HELEN LAMP	Page 1 of 7

Company / ISO Section	Criteria of ISO Section	Auditor Comments / Issues
Viamed Ltd ISO13485:2016 7.5.1	<p><b>Control of production and service provision</b></p> <p>Production and service provision shall be planned, carried out, monitored and controlled to ensure that product conforms to specification. As appropriate, production controls shall include but are not limited to:</p> <ul style="list-style-type: none"> <li>a) documentation of procedures and methods for the control of production (see 4.2.4);</li> <li>b) qualification of infrastructure;</li> <li>c) implementation of monitoring and measurement of process parameters and product characteristics;</li> <li>d) availability and use of monitoring and measuring equipment;</li> <li>e) implementation of defined operations for labelling and packaging;</li> <li>f) implementation of product release, delivery and post-delivery activities.</li> </ul> <p>The organization shall establish and maintain a record (see 4.2.5) for each medical device or batch of medical devices that provides traceability to the extent specified in 7.5.9 and identifies the amount manufactured and amount approved for distribution. The record shall be verified and approved.</p>	<p>Doc index roles + tasks management review</p>
Viamed Ltd ISO13485:2016 7.5.3	<p><b>Installation activities</b></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 organization or its supplier shall be maintained (see 4.2.5).</p>	<p>Doc index Tech files procedures</p>
Viamed Ltd ISO13485:2016 7.5.4	<p><b>Servicing activities</b></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>Doc index Tech files</p>

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	<p>The organization shall analyse records of servicing activities carried out by the organization or its supplier:</p> <ul style="list-style-type: none"> <li>a) to determine if the information is to be handled as a complaint;</li> <li>b) as appropriate, for input to the improvement process.</li> </ul> <p>Records of servicing activities carried out by the organization or its supplier shall be maintained (see 4.2.5).</p>	QA system
Viamed Ltd ISO13485:2016 7.5.6	<p><b>Validation of processes for production and service provision</b></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"> <li>a) defined criteria for review and approval of the processes;</li> <li>b) equipment qualification and qualification of personnel;</li> <li>c) use of specific methods, procedures and acceptance criteria;</li> <li>d) as appropriate, statistical techniques with rationale for sample sizes</li> <li>e) requirements for records (see 4.2.5);</li> <li>f) revalidation, including criteria for revalidation;</li> <li>g) approval of changes to the processes.</li> </ul> <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>	<p>Doc index Barcode System</p> <p>procedures Management Review</p> <p>Calibration Index</p> <p>Training Records</p>
Viamed Ltd ISO13485:2016 8.2.4	<p><b>Internal audit</b></p> <p>The organization shall conduct internal audits at planned intervals to determine whether the quality management system:</p> <ul style="list-style-type: none"> <li>a) conforms to planned and documented arrangements, requirements of this International Standard, quality management system requirements established by the organization, and applicable regulatory requirements;</li> </ul>	<p>Doc index Audit Calendar Route map</p>

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	<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. 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>Roles task.</i>
Viamed Ltd ISO13485:2016 8.4	<p><b>Analysis of data</b></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"> <li>a) feedback;</li> <li>b) conformity to product requirements;</li> <li>c) characteristics and trends of processes and product including opportunities for improvement;</li> <li>d) suppliers;</li> <li>e) audits;</li> <li>f) service reports, as appropriate.</li> </ul> <p>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.</p> <p>Records of the results of analyses shall be maintained (see 4.2.5).</p>	<i>Doc index Audit calendar QA Systems Supplier review Procedures.</i>

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

#### 1. Review Last years Audit. Update processes if required.

Are all follow on Issue resolved satisfactory.

INTRASTATS STOCK MENU – SERVICE VISITS – *Service Visits*.

All Active and Single visit services should be shown.

#### 2. Are Any service Visits Over Due

Note Tracking ID 34 in the Example,

*Nothing outstanding  
No non conformances.*

*No Service visit carried out*

Service Logs										
To Add new Servicing / Start by finding the Hospital / Company via Intrastats CRM Use the '0' Delivery account										
KEY										
Recurring Service Once Only Service										
Tracking ID	Service ID	Recur Months	Opera	Location	Description	Equipment List	Due Date	Scheduled Date	ORD's	
34	12	12	00004990	Sunderland Royal Hospital	Delivery Suite and Neonatal Unit	48 Items	26/01/15	---	ORD69051	  
33	9	12	00004260	Royal Preston Hospital	Resus cabinets, Delivery Suite	41 Items	03/04/15	---		  
35	29	12	00000780	Burnley General Hospital	Resus Cabinets and Blenders	28 Items	11/09/15	---		  
36	27	12	00000550	Royal Blackburn Hospital	Resus cabinet and blender annual service	12 Items	17/09/15	---		  
14	14	0	00005210	Walsall Manor Hospital	Tom Thumb Conversions and Upgrade	4 Items	15/02/13	---		  
13	13	0	00001350	County Durham & Darlington Hospital	maternity tom thumbs	10 Items	18/04/13	---		  
28	28	0	00000591	Royal Bolton Hospital	Resus cabinet upgrade to include blender	32 Items	17/05/13	---		  
25	18	0	00002370	Westmorland General Hospital	Resus cabinet upgrade to include blender	1 Items	19/06/14	---		  
30	23	0	00003580	North Manchester General Hosp	Resus cabinet and blender annual service	16 Items	30/07/14	---		  

Has the Over due Visit got any Action notes:

*Customer was returning items to us.*

Click:

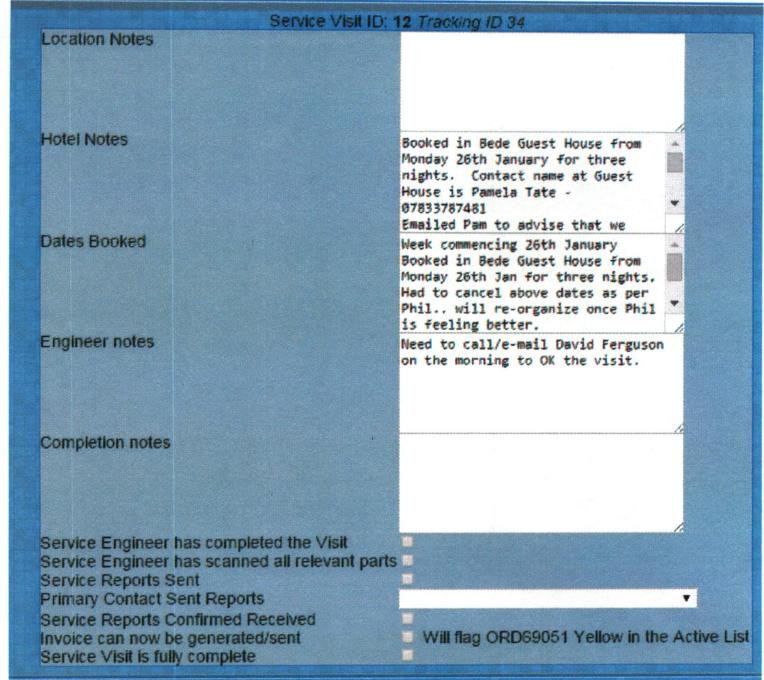


*N/A*

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

N/A

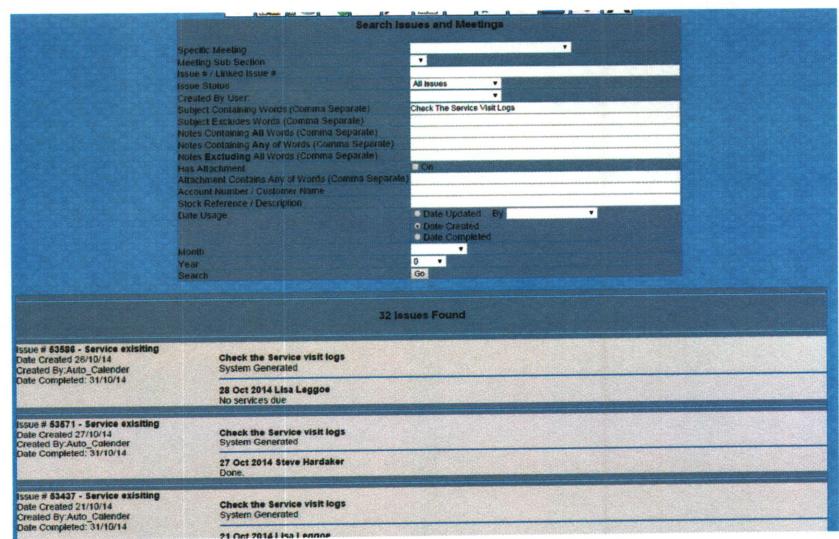
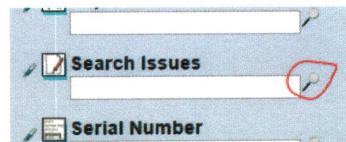


Drop out of the Service Visit Section, Search Intrastats

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

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

N/A



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**4. Are more than 1 Issue outstanding per user.**

N/A

**5. Any problems found in the Issues system not being addressed.**

No

**Sub Processes Linked to Audit**

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

**List Processes Per Title**

Clone from Docid

<b>Servicing Onsite</b>					
<b>Process Scope</b>	<b>Roll Task</b> <b>Roll Audit</b>	<b>Risk</b>	<b>Action</b>	*	Notes
PROCESSID 7985 To ensure we do not have any overdue service requirements to our customers.  Have any overdue visits not got any action notes.  NO ONSITE VISITS ANYMORE, Process not currently required	Task: 1115 357158 Company Secretary  Audit :1116 Managing Director  368505	Freq 1 Risk 1 Overall 1	Task 6M Audit 12M		
<b>Audits</b>					
<b>Process Scope</b>	<b>Roll Task</b> <b>Roll Audit</b>	<b>Risk</b>	<b>Action</b>	*	Notes

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PROCESSID <b>7889</b> To carry out Audit 24 Servicing Viamed	Task: 371427 Audit :288 <i>audit</i> Company Secretary	Freq 1 Risk 2 Overall 2	Audit 12M		
<b>Office Processes</b>					
<b>Process Scope</b>	<b>Roll Task</b> <b>Roll Audit</b>	<b>Risk</b>	<b>Action</b>	*	Notes
PROCESSID <b>5857</b> Ensuring customer onsite service visits are completed	Task: 233 372380 Office Processes <i>in terms</i> Audit :234 3709264 UK Sales Controller	Freq 2 Risk 1 Overall 2 <i>SH</i>	Task 1W Audit 1M <i>now left. more issue to someone else</i>		
PROCESSID <b>7760</b> Send letters to existing customers to remind them that a service is due on their equipment	Task: 607 372690 Marketing Processes <i>370830</i> Audit :898 Company Secretary	Freq 1 Risk 1 Overall 1	Task 1W Audit 4W		

Rolling Tasks Linked to Document :Task (233) Task (607) Task (288) Task (1115)