

## Customer Complaint and Non Conformance Review Screen

**Filter by Date Created:** From:  To:   [Show All Records \(Clear Dates\)](#)

Showing: **Viamed Issues**

[X Clear Filter](#)

- [Show All Generic Issues](#)
- [Show All Viamed Issues](#)
- [Show All Vandagraph Issues](#)
- [Show All VST Issues](#)
- [Show All Viamed Properties Issues](#)
- [Show All The Pointless Logo Company Issues](#)
- [Show All Non Minor Issues](#)
- [Next: QC 21 Active / Completed Forms](#)

Any Returns to Escalate Will Show Here :

[Show All Returns Reviews](#)

## To Filter to Company Issues you need to tag any in the Genetic Issues first

Unreviewed Telephone Complaints  
Call Call Call

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">ID2919</a>	Test Issue only so something shows up in the customer complaints review list. (based on the tick above)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Issue / Primary	Subject/Notes	Minor Internal Error i.e. not ISO	Reviewed Non Conformity	Determined Cause of Non	Evaluated action to Ensure	Planning and documenting action needed	Verify Action does not adversely	Effectiveness of corrective

ID / Call ID		Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	/ Complaint and determine if its a vigilance Issue requiring a corrective action plan	Conformity / Complaint	does not recur	and implementation QC 28b	affect Safety Performance or regulatory requirements	action reviewed
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<a href="#">ID4747</a>	test of call log complaint	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">ID5655</a>	Customer has called in response to an email he has received from Zoey saying he has an invoice overdue for £518.40 - he says that is not correct, and he should actually be in credit. Helen to look into when back from annual leave tomorrow.email back and sorted	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Non Conformance Issues

Viamed 
  Vandagraph 
  VST 
  Viamed Properties 
  The Pointless Logo Company

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No	Reviewed Non Conformity / Complaint and determine	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	if its a vigilance Issue requiring a corrective action plan					
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<a href="#">391540</a> <a href="#">11 Mar</a> <a href="#">2026</a>	Shipped Items Return to Supplier BOX1074	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
Does this Return BOX1074 warrant a NON conformance report via the CAPA process VM3COP10

Non Conformance Issues

Viamed
  Vandagraph
  VST
  Viamed Properties
  The Pointless Logo Company

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<a href="#">391539</a> <a href="#">11 Mar</a> <a href="#">2026</a>	Shipped Items Return to Supplier BOX1059	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
Does this Return BOX1059 warrant a NON conformance report via the CAPA process VM3COP10

Non Conformance Issues

Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No	Reviewed Non Conformity / Complaint and determine	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	if its a vigilance Issue requiring a corrective action plan					
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<a href="#">391298</a> <a href="#">10 Mar</a> <a href="#">2026</a>	Order Error : 162210 Viamed New Error	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Helen Lamb  
Auto Issue from Error Log 162210  
Order Entered by Catrin Hird  
Order Checked by Emily Hanson  
Office  
Error was Viamed  
New Error  
Fault:  
no stock requested to be ordered

Possible Fix  
i have requested the stock

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">391229</a> <a href="#">09 Mar</a> <a href="#">2026</a>	Order Error : 162273 Viamed Carriage - office	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Helen Lamb  
Auto Issue from Error Log 162273  
Order Entered by Kate Griffiths  
Order Checked by Aqib Majeed

Office  
 Error was Viamed  
 Carriage - office  
 Fault:  
 down as 1st class recorded but should be second class recorded, also missed by checker

Possible Fix  
 passed back to office

Audit Goods Inwards and Product Identity  
 Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">391100</a> <a href="#">09 Mar</a> <a href="#">2026</a>	Audit 09 Goods Inward And Product Identity Viamed (170)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb  
 System Generated Audit 09 BEFORE starting the Audit you need to update the Processes attached to the Audit. Find the doc in the Document Index Admin Document - Copy and Paste List Processes Per Title replacing them in the current audit.  
 Review Last years Audit see if its still suitable Carry out the Audit, reviewing and commenting on the ISO route map first. Any follow on issues must be identified with Observation Issue or a Non Conformance Issue. If the issue is a Non Conformance a QC21 form must be filled in, as per the QC21 form procedure, and investigated as such. If there are any non Conformances from the Audit, create a follow up / related Issue including all of the following - With a time for Completion Immediate Action Plan Corrective Action Plan Corrective Action Confirmation of Resolution If its a major / critical non conformance complete form QC 18  
 H.S.E. implications :No health and safety implications

Non Conformance Issues  
 Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		Failure but no requirement to Escalate Non conformance / must state in issue if its an observation						
<a href="#">390939</a> <a href="#">05 Mar 2026</a>	Order Error : 162100 Viamed Address Error - Office	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb  
Auto Issue from Error Log 162100  
Order Entered by Kate Griffiths  
Order Checked by Aqib Majeed  
Non Selected  
Error was Viamed  
Address Error - Office  
Fault:  
wrong account should have been shared services

Possible Fix

**05 Mar 2026 Helen Lamb**  
Done

Non Conformance Issues

Viamed | Vandagraph | VST | Viamed Properties | The Pointless Logo Company

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">390936</a> <a href="#">05 Mar 2026</a> 390520	Leaking Sensors Linked Issue 390520.1	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Steve Nixon  
Added by Derek Lamb sent to Steve Nixon  
FYI, looks like 0110051 we had 11 leaking sensors out of 100, its tripped a review figure due to 11 % failure rate Original Subject: 0110051 QA Data Requires Risk / Non Conformance Review Feb 2026\_ 390520.1 Added by Robert Connor sent to Derek Lamb INFORMATION ONLY ISSUE **DO NOT** ADD NOTES! QA Failures High Numbers Barcode ID Serial Number Source Status Fault / Notes 2898002 338740 Supplier Return Leaking Failed in Q.A. Did not reach end user 2898001 338739 Supplier Return Leaking Failed in Q.A. Did not reach end user 2897985 338723 Supplier Return Leaking Failed in Q.A. Did not reach end user 2897960 338698 Supplier Return Leaking. Failed in Q.A. Did not reach end user 2897963 338701 Supplier Return Leaking Failed in Q.A. Did not reach end user 2897978 338716 Supplier Return Leaking Failed in Q.A. Did

not reach end user 2898052 338790 Supplier Return Leaking Failed in Q.A. Did not reach end user 2898046 338784 Supplier Return Leaking Failed in Q.A. Did not reach end user 2898038 338776 Supplier Return Leaking Failed in Q.A. Did not reach end user 2898034 338772 Supplier Return Leaking Failed in Q.A. Did not reach end user 2898014 338752 Supplier Return Leaking Failed in Q.A. Did not reach end user

Non Conformance Issues

Viamed Vandagraph VST Viamed Properties The Pointless Logo Company

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">390872</a> <a href="#">05 Mar 2026</a>	Non conformance review history VST (285)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb

System Generated Check the history of the last Non conformance review, check actions are being carried out, and non conformances are not reoccurring. Review the Order Invoice Error Logs (link in issue) for increases in errors. Areas that are errors that are our mistake and not due to changes instigated by the customer. Note any problem areas in report. Review Customer Complaint and Non Conformance Review Screen check back over the previous month since last review to see if anything internal / staff / process related needs to be highlighted. Go to Add an issue, search non con and review any issues since the last review for internal / staff / process related issues that needs to be highlighted. Note any problems in the issue and send to Derek to review

H.S.E. implications :No health and safety implications

**06 Mar 2026 Helen Lamb**

Checked back through all Non Conformance issues (not automatically generated) to 1st March 26. Nothing Of concern or to be investigated or reported on in Non Con issues. Review VST Product Feedback Negative (742) nothing new. Review VST Product Feedback Positive (1190) nothing new. Review VST Feedback - Customer Feedback Negative (740) nothing new. Review VST Customer Feedback Positive(1191) nothing new. Review VST Feedback - Customer Complaints (738) nothing new. Order Invoice Error Logs - no issues, nothing to worry about. Non Conformance, complaints and feedback headers nothing new, no concerns. Non Conformance issues review screen - No issues not already reviewed above in log. Only issues are those reviewing meeting headers from the Management Review. Nothing raised. No other feedback issues relating to non conformances that need to be monitored or reviewed. Derek please review

**06 Mar 2026 Derek Lamb**

thankyou

Non Conformance Issues

Viamed Vandagraph VST Viamed Properties The Pointless Logo Company

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		no requirement to Escalate Non conformance / must state in issue if its an observation						
<a href="#">390577</a> <a href="#">03 Mar 2026</a>	Documentation out of date (372)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb  
System Generated  
Check for Out of Date documents

This is an audit, you do not need to perform the Task:

Simply ensure all out of date documents have an Issue attached to get them updated.  
If the Issue is more than 2 Months out of date read the issue - if appropriate generate a non conformance Issue

ISO - Document index admin

Scroll down and check if any documents have gone out of date,

Either update the document or create an Issue to the relevant person from the document admin / details screen.  
Remember if you update a document reset the expiry date  
H.S.E. implications :No health and safety implications  
**03 Mar 2026 Helen Lamb**  
Done

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">390521</a> <a href="#">02 Mar 2026</a>	0110127 QA Data Requires Risk / Non Conformance Review Feb 2026	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
Added by Robert Connor sent to Derek Lamb  
INFORMATION ONLY ISSUE **DO NOT** ADD NOTES!  
QA Failures High Numbers  
**04 Mar 2026 Derek Lamb**  
automotive sensor, known yield  
**04 Mar 2026 Robert Connor**

Done

Non Conformance Issues  
Viamed

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<a href="#">390520</a> <a href="#">02 Mar 2026</a>	0110051 QA Data Requires Risk / Non Conformance Review Feb 2026	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
Added by Robert Connor sent to Derek Lamb  
INFORMATION ONLY ISSUE **DO NOT** ADD NOTES!  
QA Failures High Numbers

**05 Mar 2026 Derek Lamb**  
Created Related Issue #390936  
Added by Derek Lamb sent to Steve Nixon  
FYI, looks like 0110051 we had 11 leaking sensors out of 100, its tripped a review figure due to 11 % failure rate Original Subject: 0110051 QA Data Requires Risk / Non Conformance Review Feb 2026\_390520.1 Added by Robert Connor sent to Derek Lamb INFORMATION ONLY ISSUE **DO NOT** ADD NOTES! QA Failures High Numbers Barcode ID Serial Number Source Status Fault / Notes 2898002 338740 Supplier Return Leaking Failed in Q.A. Did not reach end user 2898001 338739 Supplier Return Leaking Failed in Q.A. Did not reach end user 2897985 338723 Supplier Return Leaking Failed in Q.A. Did not reach end user 2897960 338698 Supplier Return Leaking. Failed in Q.A. Did not reach end user 2897963 338701 Supplier Return Leaking Failed in Q.A. Did not reach end user 2897978 338716 Supplier Return Leaking Failed in Q.A. Did not reach end user 2898052 338790 Supplier Return Leaking Failed in Q.A. Did not reach end user 2898046 338784 Supplier Return Leaking Failed in Q.A. Did not reach end user 2898038 338776 Supplier Return Leaking Failed in Q.A. Did not reach end user 2898034 338772 Supplier Return Leaking Failed in Q.A. Did not reach end user 2898014 338752 Supplier Return Leaking Failed in Q.A. Did not reach end user

**06 Mar 2026 Derek Lamb**  
informed SN to see if we have a problem

**09 Mar 2026 Robert Connor**  
Done

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	requiring a corrective action plan						
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<a href="#">390519</a> <a href="#">02 Mar 2026</a>	0110137 QA Data Requires Risk / Non Conformance Review Feb 2026	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
 Added by Robert Connor sent to Derek Lamb  
 INFORMATION ONLY ISSUE **DO NOT** ADD NOTES!  
 QA Failures High Numbers  
**04 Mar 2026 Derek Lamb**  
 automotive sensor, known yield  
**04 Mar 2026 Robert Connor**  
 Done

Viamed Audits Calander  
 Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">390302</a> <a href="#">27 Feb 2026</a>	Audits Calander - Audits Calander	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
 Added by Derek Lamb sent to Derek Lamb  
 the newer audits need adding to the audits calender so i can close them off

Non Conformance Issues  
 Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non	Reviewed Non Conformity /	Determined Cause of Non	Evaluated action to Ensure	Planning and documenting action needed and	Verify Action does not adversely affect Safety	Effectiveness of corrective action reviewed
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		Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Complaint and determine if its a vigilance Issue requiring a corrective action plan	Conformity / Complaint	does not recur	implementation QC 28b	Performance or regulatory requirements	
<a href="#">389899</a> <a href="#">24 Feb 2026</a> 380348	BOX1054 Linked Issue 380348.1	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
 Added by Derek Lamb sent to Derek Lamb  
 Original Subject: Shipped Items Return to Supplier BOX1054\_380348.1 Leaking gas ? the figure seems unusually high. We?re waiting for a response from the supplier. issue untagged still yellow but low do the list

Non Conformance Issues  
 Viamed

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<a href="#">389811</a> <a href="#">23 Feb 2026</a>	Shipped Items Return to Supplier BOX1063	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
 Does this Return BOX1063 warrant a NON conformance report via the CAPA process VM3COP10  
**02 Mar 2026 Derek Lamb**  
 no capa required, Envitec Oxygen monitor MySign O going back to manufacturer for review

Non Conformance Issues  
 Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">389517</a> <a href="#">19 Feb 2026</a>	Order Error : 161533 Viamed Carriage - office	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Helen Lamb  
Auto Issue from Error Log 161533  
Order Entered by Aqib Majeed  
Order Checked by Emily Hanson  
Office  
Error was Viamed  
Carriage - office  
Fault:  
spelling mistake on carriage line, missed by imputter and checker.

Possible Fix  
changed at time of picking

**20 Feb 2026 Helen Lamb**  
Done

Audit Contract Review

Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	action plan						
<a href="#">388821</a> <a href="#">13 Feb 2026</a>	Shipped Items Return to Supplier BOX1069	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
Does this Return BOX1069 warrant a NON conformance report via the CAPA process VM3COP10

**16 Feb 2026 Derek Lamb**  
unit going back to bp for repair, originally sold 2013

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">388682</a> <a href="#">11 Feb 2026</a>	Order Error : 161745 Vandagraph Carriage - office	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Helen Lamb  
Auto Issue from Error Log 161745  
Order Entered by Kate Griffiths  
Order Checked by Sophie Lines  
Office  
Error was Vandagraph Carriage - office  
Fault:  
Used royal mail for northern ireland even though there is a delivery helper tagged to the order page that states that northern ireland should be UPS

Possible Fix  
Passed back to the office

**12 Feb 2026 Helen Lamb**  
Done



		to Escalate Non conformance / must state in issue if its an observation						
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<a href="#">388487</a> <a href="#">10 Feb 2026</a>	Order Error : 161409 Viamed Account error	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Helen Lamb  
Auto Issue from Error Log 161409  
Order Entered by Aqib Majeed  
Order Checked by Kate Griffiths  
Office  
Error was Viamed  
Account error  
Fault:  
shouldnt have been charged vat. checking missed it as well

Possible Fix

**12 Feb 2026 Helen Lamb**  
Done  
**12 Feb 2026 Helen Lamb**  
Done

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">388465</a> <a href="#">10 Feb 2026</a>	Order Error : 161612 Vandagraph New Error	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Helen Lamb  
Auto Issue from Error Log 161612  
Order Entered by Kate Griffiths  
Order Checked by Sophie Lines  
Non Selected  
Error was Vandagraph  
New Error  
Fault:  
Order not fulfilled on Shopify

Possible Fix

Done

10 Feb 2026 Helen Lamb

Done

Audits Meeting Closure

Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">388321</a> <a href="#">09 Feb 2026</a>	Audit 27 Software Validation Viamed (821)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb

System Generated Task To be Completed Audit 27 Software Validation To confirm the Prime functions of the Software used is verified. The Audit itself, Intrastats, physical process being carried out. Complete Audit 27 BEFORE starting the Audit you need to update the Processes attached to the Audit. Find the doc in the Document Index Admin Document - Copy and Paste List Processes Per Title replacing them in the current audit. Review Last years Audit see if its still suitable Carry out the Audit, reviewing and commenting on the ISO route map first. Any follow on issues must be identified with Observation Issue or a Non Conformance Issue. If the issue is a Non Conformance a QC21 form must be filled in, as per the QC21 form procedure, and investigated as such. If there are any non Conformances from the Audit, create a follow up / related Issue including all of the following - With a time for Completion Immediate Action Plan Corrective Action Plan Corrective Action Confirmation of Resolution If its a major / critical non conformance complete form QC 18  
H.S.E. implications :No health and safety implications

Audits Closure

Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		Non conformance / must state in issue if its an observation						
<a href="#">388026</a> <a href="#">05 Feb 2026</a>	Objective To agree the current schedule of Top Level audits.	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

General  
05 Feb 2026 Derek Lamb  
All agree to current schedule of top level audits.  
All agreed with 2025 audits results. Closed 2025 audits.  
All Agree to the 2026 top level audits, Audit schedule .

Results of internal audits / Mini Audits  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">388024</a> <a href="#">05 Feb 2026</a>	Objective Review the results of internal audits / Mini Audits over the last 12 months. Check everyone is happy with the status of the current years Audits. Reviewed Audit scheduled for next year.	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

General  
05 Feb 2026 Derek Lamb  
Reviewed Internal Paper Main Audits 2025 all happy, Reviewed Mini Audit Management Reviews Global all happy, Reviewed Mini Audit Reviews outstanding Issues, Reviewed VST Audit Calendar 2026, Discussed follow up issues to 2025 Main Audits.  
ISO 9001:2015+A1:2024(E) is Current as per our certification.  
All agreed happy with current years Audits.  
Reviewed all Audits carried out.  
All agreed to continue with existing Audit calendar and scheduled for 2026.  
Next BSI Audit will be February 2026.  
We reviewed all reports live on the system, nothing flagged up in 2025 audits.

Non Conformities Review  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">387975</a> <a href="#">05 Feb 2026</a>	Objective To review and action on any non conformances within 15 days. Check for new qc 21 forms.	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

General  
05 Feb 2026 Derek Lamb  
reviewed non conformance headers, issues sections, contain audits, and supplier returns, all supplier returns reviewed to see if a capa should be raised, all normal sensor fail types, so did not require a capa/qc21 form

Non Conformance Issues Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">387858</a> <a href="#">05 Feb 2026</a>	Non conformance review history Viamed (283)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb  
System Generated Check the history of the last Non conformance review, check actions are being carried out, and non

conformances are not reoccurring. Review the Order Invoice Error Logs (link in issue) for increases in errors. Areas that are errors that are our mistake and not due to changes instigated by the customer. Note any problem areas in report. Review Customer Complaint and Non Conformance Review Screen check back over the previous month since last review to see if anything internal / staff / process related needs to be highlighted. Go to Add an issue, search non con and review any issues since the last review for internal / staff / process related issues that needs to be highlighted. Note any problems in the issue and send to Derek to review

H.S.E. implications :No health and safety implications

**20 Feb 2026 Helen Lamb**

Checked back through all Non Conformance issues (not automatically generated) to 1st February 26. 384994 0110361 QA Data Requires Risk. Nothing of concern or to be investigated or reported on in Non Con issues. Review VIAMED Feedback - Product Feedback Negative (741) nothing new. Review VIAMED Feedback - Product Feedback Positive (1189) nothing new. Review VIAMED Feedback - Customer Feedback Negative (739) nothing new. Review VIAMED Feedback - Customer Feedback Positive (1188) nothing new. Review VIAMED Feedback - Customer Complaints (737) nothing new. Order Invoice Error Logs - Only minor issue, nothing to worry about. Non Conformance, complaints and feedback headers nothing new no concerns. We have new issues relating to Bluepoint asking us for feedback. Nothing of concern. Nothing not already reviewed in the non conformance log all just general working errors. Nothing to worry about and nothing excessive. Nothing else new not already covered. Non Conformance issues review screen - No issues not already reviewed above in log. No other feedback issues relating to non conformances that need to be monitored or reviewed. Derek please review the above Done

**Audit Analysis of Data**

Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">387853</a> <a href="#">05 Feb 2026</a>	Review Company Data (114)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb

System Generated Review Sales - Number Invoices / Credits per day reports Search for any potential problems in the Data provided

Employee Menu Check Audit Roles Titles and Processes in Employee Roles and Titles. Check the Page for Red Crosses and potentially missing objectives Search the Outstanding Jobs column for potential problems Do we have enough resources. Check the Issues Statistics in the meeting header, Check Issues are being completed in a timely manner.

H.S.E. implications :No health and safety implications

**05 Feb 2026 Derek Lamb**

OverView :05/02/2026 SectionDate UpdatedValue Viamed Forward Orders02/02/2026£ 128,711.00 Viamed Back Orders02/02/2026£ 143,080.70 Viamed Bank Balance02/02/2026£ 227,663.35 Viamed Debtors02/02/2026£ 225,157.62 Viamed Creditors02/02/2026£ 211,813.49 Barclays Loan to Viamed:02/02/2026£ 0.00 Vandagraph Loan To Viamed:02/02/2026£ 0.00 VST Loan To Viamed:02/02/2026£ 0.00 Viamed Properties Loan to Viamed :£0.00 Loan from GGL JSL to Viamed:02/02/2026£ 0.00 Loan from DL To Viamed:02/02/2026£ 0.00 Loan from SN to Viamed:02/02/2026£ 0.00 VST Bank Balance£ 60691.38 Vandagraph Bank Balance£ 155136.09 Viamed Stock Valuation05/02/2026£ 0 Viamed Turnover30/01/26£ 246,056.59 Vst Turnover30/01/26£ 82,321.07 Vand Turnover30/01/26£ 26,473.50 VST Debtors£ 51315 VST Creditors£ 54846 VST Back Orders£ 33149 VST Forward Orders£ 419960 Vandagraph Debtors£ 15964 Vandagraph Creditors£ 15693 Vandagraph Back Orders£ 3368 Vandagraph Forward Orders£ 0

**Non Conformance Issues**

Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">387730</a> <a href="#">04 Feb 2026</a>	Non conformance review history Viamed (284)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
 System Generated Check the below review is being carried out Check the history of the last Non conformance review, check actions are being carried out, and non conformances are not reoccurring  
 H.S.E. implications :No health and safety implications  
**06 Feb 2026 Derek Lamb**  
 task is upto date just the current one due

Non Conformance Issues  
 Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">387666</a> <a href="#">03 Feb 2026</a>	Order Error : 161377 Viamed Date error	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb  
 Auto Issue from Error Log 161377  
 Order Entered by Sophie Lines  
 Order Checked by Kate Griffiths

Office

Error was Viamed

Date error

Fault:

Customer order states delivery in June, memo on order states delivery in June, order been placed as delivery in January.

Possible Fix

Edited due date to 1st June.

**03 Feb 2026 Helen Lamb**

Done

Non Conformance Issues

Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">386992</a> <a href="#">26 Jan 2026</a>	Shipped Items Return to Supplier BOX1064	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb

Does this Return BOX1064 warrant a NON conformance report via the CAPA process VM3COP10

**28 Jan 2026 Derek Lamb**

no issue just a calibration

Audit Organisation and Process Verification Internal Process Verification

Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		to Escalate Non conformance / must state in issue if its an observation						
<a href="#">386202</a> <a href="#">19 Jan 2026</a>	Email Distribution (366)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
System Generated Check the global mailbox, there should be no mails more than a Day old. Give allowance for Monday mornings, there should be no emails older than Friday afternoon. Also note size of space used in Global inbox H.S.E. implications :No health and safety implications  
**19 Jan 2026 Derek Lamb**  
inbox empty 10am monday

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">385909</a> <a href="#">14 Jan 2026</a>	Order Error : 161110 Viamed Memo error	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Helen Lamb  
Auto Issue from Error Log 161110  
Order Entered by Aqib Majeed  
Order Checked by Kate Griffiths  
Office  
Error was Viamed  
Memo error  
Fault:  
Did not state on the order/ memo that these sensors were to be unboxed, only realised after picking and packing the order that the box size stated on the order was too small, and checked it and the office were given two different box sizes for boxed and unboxed, email was attached to the order to state they will try unboxed.  
Possible Fix  
**20 Jan 2026 Helen Lamb**  
staff involved were spoken to at the time and are aware of the mistake.

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">385451</a> <a href="#">08 Jan 2026</a>	Order Error : 161012 Vandagraph Spelling Mistake - office	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb

Auto Issue from Error Log 161012

Order Entered by Kate Griffiths

Order Checked by Aqib Majeed

Office

Error was Vandagraph

Spelling Mistake - office

Fault:

surname spelt two different ways on the order, been copied from web page but looks like no one has contact the customer to check which spelling is correct.

Possible Fix

passed back to office

**09 Jan 2026 Helen Lamb**

Done

Non Conformance Issues

Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		/ must state in issue if its an observation						
<a href="#">385109</a> <a href="#">06 Jan 2026</a>	Order Error : 160943 Viamed New Error	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb  
Auto Issue from Error Log 160943  
Order Entered by Kate Griffiths  
Order Checked by Aqib Majeed  
Goods Out  
Error was Viamed  
New Error  
Fault:  
Order not fulfilled on Shopify

Possible Fix  
Done

**06 Jan 2026 Helen Lamb**  
Done  
**06 Jan 2026 Helen Lamb**  
Done

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">384994</a> <a href="#">05 Jan 2026</a>	0110361 QA Data Requires Risk / Non Conformance Review Dec 2025	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
Added by Robert Connor sent to Derek Lamb  
INFORMATION ONLY ISSUE **DO NOT** ADD NOTES!  
QA Failures High Numbers  
**06 Jan 2026 Derek Lamb**  
0110361Oxygen sensor R-22AHJR Jikco OEM sensorExamine 7 / 42 16.67%Examine 10 / 107 9.35% / 10 /Examine 34 / 277 12.27%Examine 12 / 133 9.02% / 0 / 0 / 10 / 0 /Examine 14 / 100 14.00% jiko sensor, known yield, no cap required  
**06 Jan 2026 Robert Connor**  
Done

Non Conformance Issues

Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">384914</a> <a href="#">05 Jan 2026</a>	Non conformance review history Viamed (283)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb

System Generated Check the history of the last Non conformance review, check actions are being carried out, and non conformances are not reoccurring. Review the Order Invoice Error Logs (link in issue) for increases in errors. Areas that are errors that are our mistake and not due to changes instigated by the customer. Note any problem areas in report. Review Customer Complaint and Non Conformance Review Screen check back over the previous month since last review to see if anything internal / staff / process related needs to be highlighted. Go to Add an issue, search non con and review any issues since the last review for internal / staff / process related issues that needs to be highlighted. Note any problems in the issue and send to Derek to review

H.S.E. implications :No health and safety implications

**09 Jan 2026 Helen Lamb**

Checked back through all Non Conformance issues (not automatically generated) to 1st January 26. Nothing of concern or to be investigated or reported on in Non Con issues. Review VIAMED Feedback - Product Feedback Negative (741) nothing new. Review VIAMED Feedback - Product Feedback Positive (1189) nothing new. Review VIAMED Feedback - Customer Feedback Negative (739) nothing new. Review VIAMED Feedback - Customer Feedback Positive (1188) nothing new. Review VIAMED Feedback - Customer Complaints (737) nothing new. Order Invoice Error Logs - only one data entry error no other issue, nothing to worry about. Non Conformance, complaints and feedback headers nothing new no concerns. Nothing not already reviewed in the non conformance log all just general working errors. Nothing to worry about and nothing excessive. Nothing else new not already covered. Non Conformance issues review screen - No issues not already reviewed above in log. No other feedback issues relating to non conformances that need to be monitored or reviewed. Derek please review the above

**12 Jan 2026 Derek Lamb**

thankyou

Audit Analysis of Data

Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		no requirement to Escalate Non conformance / must state in issue if its an observation						
<a href="#">384905</a> <a href="#">05 Jan 2026</a>	Review Company Data (114)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
System Generated Review Sales - Number Invoices / Credits per day reports Search for any potential problems in the Data provided

Employee Menu Check Audit Roles Titles and Processes in Employee Roles and Titles. Check the Page for Red Crosses and potentially missing objectives Search the Outstanding Jobs column for potential problems Do we have enough resources. Check the Issues Statistics in the meeting header, Check Issues are being completed in a timely manner.  
H.S.E. implications :No health and safety implications  
**06 Jan 2026 Derek Lamb**  
Done

**Audit Picking and Packing Viamed**

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">384634</a> <a href="#">01 Jan 2026</a>	Audit 01 Picking Packing Viamed (24)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
System Generated BEFORE starting Audit you need to update the Processes attached to the Audit. Find the doc in the Document Index Admin Document - Copy and Paste List Processes Per Title replacing them in the current audit. Review Last years Audit see if its still suitable Carry out the Audit, reviewing and commenting on the ISO route map first. Any follow on issues must be identified with Observation Issue or a Non Conformance Issue. If the issue is a Non Conformance a QC21 form must be filled in, as per the QC21 form procedure, and investigated as such. If there are any non Conformances from the Audit, create a follow up / related Issue including all of the following - With a time for Completion Immediate Action Plan Corrective Action Plan Corrective Action Confirmation of Resolution If its a major / critical non conformance complete form QC 18  
H.S.E. implications :No health and safety implications  
**14 Jan 2026 Helen Lamb**  
audit done, nothing outstanding no non conformances. No problems have been found this area has had a lot of programming to make it easier and less prone to mistakes.  
**21 Jan 2026 Derek Lamb**  
thankyou

Audit Design Control  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">384633</a> <a href="#">01 Jan 2026</a>	Audit 03 Design Control Viamed (22)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
 Audit 03 Design Control Review Last years Audit see if its still suitable Before Proceeding you need to update the Processes attached to the Audit Search the Document in the Index, View the Admin Page Copy and Paste the Attached Processes, replacing them in the current audit Any follow on issues must be identified with Observation Issue or a Non Conformance Issue. If the issue is a Non Conformance a QC21 form must be filled in, as per the QC21 form procedure, and investigated as such. Any non Conformances from the Audit: Create a follow up / related Issue, With a time for Completion Immediate Action Plan Corrective Action Plan Corrective Action Confirmation of Resolution if its a major / critical non conformance complete form QC 18  
 H.S.E. implications :No health and safety implications  
**19 Feb 2026 Helen Lamb**  
 Audit completed no issues either observations or non conformances. Please review this  
**20 Feb 2026 Derek Lamb**  
 thankyou

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		an observation						
<a href="#">384513</a> <a href="#">30 Dec 2025</a>	Order Error : 160911 Viamed Carriage - office	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb  
Auto Issue from Error Log 160911  
Order Entered by Aqib Majeed  
Order Checked by Michael Lamb  
Office  
Error was Viamed  
Carriage - office  
Fault:  
Down as free carriage, but this customer is a distributor and should pay for carriage.

Possible Fix  
passed back to office for correction

**30 Dec 2025 Helen Lamb**  
Done

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">383890</a> <a href="#">18 Dec 2025</a>	eBay Order Should be VAT Exempt	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Helen Lamb  
Added by Catrin Hird sent to Helen Lamb  
Please can you amend CVM160748 as it should have been VAT exempt. Proof of export attached to order. eBay order 17-13968-62623  
**29 Dec 2025 Helen Lamb**  
Done  
**30 Dec 2025 Catrin Hird**  
Sorry, VAT had been charged and shows on invoice against order, how do I see if there has been a credit put against the order? Apologies if I've misunderstood :)  
**06 Jan 2026 Helen Lamb**  
sorry i corrected the involve, didnt do a credit and fixed invoice in Xero  
**08 Jan 2026 Catrin Hird**  
Presume all ok now Done

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">383889</a> <a href="#">18 Dec 2025</a>	Order Error : 160748 Viamed Ebay error - office	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Helen Lamb  
Auto Issue from Error Log 160748  
Order Entered by Sophie Lines  
Order Checked by Kate Griffiths  
Office  
Error was Viamed  
Ebay error - office  
Fault:  
Order should have been VAT exempt, order was being exported straight out of the country. Proof of export attached to documentation.

Possible Fix  
Not charge VAT on Global Shipping Service orders.

**19 Dec 2025 Helen Lamb**  
Done

Audit Organisation and Process Verification Internal Process Verification  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		in issue if its an observation						
<a href="#">383699</a> <a href="#">17 Dec 2025</a>	Email Distribution (366)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
System Generated Check the global mailbox, there should be no mails more than a Day old. Give allowance for Monday mornings, there should be no emails older than Friday afternoon. Also note size of space used in Global inbox  
H.S.E. implications :No health and safety implications  
**23 Dec 2025 Derek Lamb**  
Email box empty and upto date

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">383494</a> <a href="#">15 Dec 2025</a>	Order Error : 160197 Viamed Missing info - office	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Helen Lamb  
Auto Issue from Error Log 160197  
Order Entered by Aqib Majeed  
Order Checked by Kate Griffiths  
Office  
Error was Viamed  
Missing info - office  
Fault:  
Missed wording stated on order that they require a test certificate, also missed by checker. Customer has asked for one, but we are unable to now as the unit is now with them. The request was made on the PO right underneath the stock that they requested  
Possible Fix  
  
**15 Dec 2025 Helen Lamb**  
Done

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non	Reviewed Non Conformity /	Determined Cause of Non	Evaluated action to Ensure	Planning and documenting action needed and	Verify Action does not adversely affect Safety	Effectiveness of corrective action reviewed
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2025

Derek Lamb

Added by Robert Connor sent to Helen Lamb  
INFORMATION ONLY ISSUE **DO NOT** ADD NOTES!

We have had 4 instances in the last month of office staff not taking action on stock memos which has delayed the shipping of orders that otherwise could have been sent out very quickly. We have several part numbers which are not kept in regular stock as they sell infrequently, but can be made up from regular stock as-needed, these stock items all have memos instructing the office to inform the warehouse when putting an order on for these items. Part numbers for these are 0110073 R22VA matched pair, and 0021015 case of 96 Posey wraps. We have had multiple orders in the last month where customer orders for these items have been placed without informing me that they need to be made up, resulting in customer orders sitting unfulfilled for up to 3 weeks in one case. The most recent example with 0021015s was placed after I had already reminded the office that they must inform the warehouse about this kind of order. It is very important that people read and take action on stock memos when they are placing orders as they are generally added for good reason.

**06 Jan 2026 Helen Lamb**

spoke to the office about this but also Derek said he is looking at putting stock memo on the order entry page. Hopefully both will help changing user to Derek as a nudge

**06 Jan 2026 Helen Lamb**

spoke to the office about this but also Derek said he is looking at putting stock memo on the order entry page. Hopefully both will help changing user to Derek as a nudge

**06 Jan 2026 Derek Lamb**

ok first implementation of stock memos on order entry screen has been done.

**06 Jan 2026 Robert Connor**

Sounds good thanks, I'll try too keep an eye on this and see how it turns out.

Non Conformance Issues

Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">382670</a> <a href="#">05 Dec 2025</a>	Non conformance review history Viamed (283)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb

System Generated Check the history of the last Non conformance review, check actions are being carried out, and non conformances are not reoccurring. Review the Order Invoice Error Logs (link in issue) for increases in errors. Areas that are errors that are our mistake and not due to changes instigated by the customer. Note any problem areas in report. Review Customer Complaint and Non Conformance Review Screen check back over the previous month since last review to see if anything internal / staff / process related needs to be highlighted. Go to Add an issue, search non con and review any issues since the last review for internal / staff / process related issues that needs to be highlighted. Note any problems in the issue and send to Derek to review

H.S.E. implications :No health and safety implications

**22 Dec 2025 Helen Lamb**

Checked back through all Non Conformance issues (not automatically generated) to 1st December 25. Nothing of concern or to be investigated or reported on in Non Con issues. Review VIAMED Feedback - Product Feedback Negative (741) nothing new. Review VIAMED Feedback - Product Feedback Positive (1189) nothing new. Review VIAMED Feedback - Customer Feedback Negative (739) nothing new. Review VIAMED Feedback - Customer Feedback Positive (1188) nothing new. Review VIAMED Feedback - Customer Complaints (737) nothing new. Order Invoice Error Logs - only one data entry error no other issue, nothing to worry about. Non Conformance, complaints and feedback headers 381787 Honeywell error shipping wrong

senor in box, issues 381351, 381375, 381653, 381760, 381892, 381958, 381987, 382119, 382537, 382613, 382624, 382914, 382953, 383214, 383215, 383216, 383217, 383494, 383637, 383889, 383890. These are all office errors from the error log. We have staff ill and this has affected the amount of errors we have been having. Will monitor these in the new year and see if we need to do anything to make this more efficient. Nothing else new not already covered. Non Conformance issues review screen - No issues not already reviewed above in log. No other feedback issues relating to non conformances that need to be monitored or reviewed. Derek please review the above Done

Audit Analysis of Data  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">382664</a> <a href="#">05 Dec</a> <a href="#">2025</a>	Review Company Data (114)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb

System Generated Review Sales - Number Invoices / Credits per day reports Search for any potential problems in the Data provided

Employee Menu Check Audit Roles Titles and Processes in Employee Roles and Titles. Check the Page for Red Crosses and potentially missing objectives Search the Outstanding Jobs column for potential problems Do we have enough resources. Check the Issues Statistics in the meeting header, Check Issues are being completed in a timely manner.

H.S.E. implications :No health and safety implications

**11 Dec 2025 Derek Lamb**

reviewed Viamed Forward Orders 02/12/2025£ 81,439.00 Viamed Back Orders 02/12/2025£ 160,393.66 Viamed Bank Balance 02/12/2025£ 169,260.01 Viamed Debtors 02/12/2025£ 230,285.49 Viamed Creditors 02/12/2025£ 203,688.41

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		to Escalate Non conformance / must state in issue if its an observation						
<a href="#">382613</a> <a href="#">04 Dec</a> <a href="#">2025</a>	Order Error : 160171 Viamed New Error	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb  
Auto Issue from Error Log 160171  
Order Entered by Kate Griffiths  
Order Checked by Emily Hanson  
Office  
Error was Viamed  
New Error  
Fault:  
down as CIP but is EXW ..NO courier details EC 3/12/25

Possible Fix

**05 Dec 2025 Helen Lamb**  
Done

Audit Internal Audits  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">382126</a> <a href="#">01 Dec</a> <a href="#">2025</a>	Audit 17 Internal Audits VST (191)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
System Generated Audit 17 BEFORE starting the Audit you need to update the Processes attached to the Audit. Find the doc in the Document Index Admin Document - Copy and Paste List Processes Per Title replacing them in the current audit.  
Review Last years Audit see if its still suitable Carry out the Audit, reviewing and commenting on the ISO route map first. Any follow on issues must be identified with Observation Issue or a Non Conformance Issue. If the issue is a Non Conformance a QC21 form must be filled in, as per the QC21 form procedure, and investigated as such. If there are any non Conformances from the Audit, create a follow up / related Issue including all of the following - With a time for Completion Immediate Action Plan Corrective Action Plan Corrective Action Confirmation of Resolution If its a major / critical non conformance complete form QC 18  
H.S.E. implications :No health and safety implications  
**31 Dec 2025 Helen Lamb**  
completed audit attached please review. Nothing outstanding no non conformances.  
**31 Dec 2025 Derek Lamb**

Done  
**31 Dec 2025 Derek Lamb**  
done

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">382119</a> <a href="#">28 Nov 2025</a>	Order Error : 160173 Viamed New Error	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb  
Auto Issue from Error Log 160173  
Order Entered by Kate Griffiths  
Order Checked by Aqib Majeed  
Goods Out  
Error was Viamed  
New Error  
Fault:  
Order not fulfilled on ebay

Possible Fix  
Done

**30 Nov 2025 Helen Lamb**  
Done

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		no requirement to Escalate Non conformance / must state in issue if its an observation						
<a href="#">381987</a> <a href="#">27 Nov 2025</a>	Order Error : 156530 Viamed Checking error	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb  
Auto Issue from Error Log 156530  
Order Entered by Emily Hanson  
Order Checked by Sophie Lines  
Office  
Error was Viamed  
Checking error  
Fault:  
Order put on the wrong account. Should have been Victoria Blackpool not Scotland.

Possible Fix

**28 Nov 2025 Helen Lamb**  
duplicate issue Done

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">381986</a> <a href="#">27 Nov 2025</a>	Order Error : 156530 Viamed Address Error - Office	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb  
Auto Issue from Error Log 156530  
Order Entered by Emily Hanson  
Order Checked by Sophie Lines  
Office  
Error was Viamed  
Address Error - Office  
Fault:  
Order put on the wrong account. Should have been Victoria Blackpool not Scotland.

Possible Fix

**28 Nov 2025 Helen Lamb**  
duplicate issue Done

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">381985</a> <a href="#">27 Nov 2025</a>	Order Error : 156530 Viamed Address Error - Office	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb  
Auto Issue from Error Log 156530  
Order Entered by Emily Hanson  
Order Checked by Sophie Lines  
Office  
Error was Viamed  
Address Error - Office  
Fault:  
Order put on the wrong account. Should have been Victoria Blackpool not Scotland.  
  
Possible Fix

**28 Nov 2025 Helen Lamb**  
Done

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">381892</a> <a href="#">26 Nov 2025</a>	PVM3772 Due Nov 2024, not chased or delivered	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
Added by Catrin Hird sent to Derek Lamb  
PVM3772 Due Nov 2024, not chased or delivered Customer now placed an order.  
**01 Dec 2025 Derek Lamb**  
Supplier Issue, order should be on its way now,  
**09 Dec 2025 Catrin Hird**  
Done

Supplier Complaints (us to them)  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">381787</a> <a href="#">25 Nov 2025</a>	Honeywell error shipping wrong sensor in box	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Catherine Spence  
Added by Helen Lamb sent to Catherine Spence  
Honeywell have sold us a R-47 0110047 but the sensor inside is 0110814 OOMLF204. The outer pack says R-47 0110047 V107762 expire date 2025-03-01. The sensor inside says OOMLF204 B110191 expire date 2024-04-01 Photos attached. Its

definitely the wrong sensor in side the customer has been asked to return this to us and we will be sending a FOC replacement out to them now. Cathy to let me know when this comes back in.

**16 Dec 2025 Catherine Spence**

noted

**29 Jan 2026 Catherine Spence**

Done

Non Conformance Issues

Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">381653</a> <a href="#">24 Nov 2025</a>	Order Error : 160261 Vandagraph Carriage - office	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb

Auto Issue from Error Log 160261

Order Entered by Sophie Lines

Order Checked by Emily Hanson

Office

Error was Vandagraph

Carriage - office

Fault:

Wrong priority on the order

Possible Fix

sent back to office for correction

**24 Nov 2025 Helen Lamb**

Done

Non Conformance Issues

Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	action plan					
<a href="#">381375</a> <a href="#">20 Nov 2025</a>	Order Error : 158795 Viamed Missing info - office	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb  
Auto Issue from Error Log 158795  
Order Entered by Kate Griffiths  
Order Checked by Sophie Lines  
Non Selected  
Error was Viamed  
Missing info - office  
Fault:  
requirement on PO was not mentioned on the order and also missed by the checker

Possible Fix  
passed back to the office to correct

**24 Nov 2025 Helen Lamb**  
Done

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">381351</a> <a href="#">20 Nov 2025</a>	Order Error : 160236 Viamed Missing info - office	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb  
Auto Issue from Error Log 160236  
Order Entered by Aqib Majeed  
Order Checked by Sophie Lines  
Office  
Error was Viamed  
Missing info - office  
Fault:

email address for contact was entered as the PO number,

Possible Fix  
sent back to office to correct

**20 Nov 2025 Helen Lamb**  
Done

**Audit Documentation Control**  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">381136</a> <a href="#">18 Nov 2025</a>	PAQ required: 0111261	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Catrin Hird  
Added by Catrin Hird sent to Catrin Hird  
for QVM159991 & QVM160136  
**19 Nov 2025 Catrin Hird**  
Updated name and date. Need information from Maxtec to change to new versions.

**Audit Contract Review**  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	action plan					
<a href="#">380933</a> <a href="#">14 Nov 2025</a>	Shipped Items Return to Supplier BOX1056	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
Does this Return BOX1056 warrant a NON conformance report via the CAPA process VM3COP10

**14 Nov 2025 Derek Lamb**  
the leaky sensors going back to supplier already mention in other reviews

Audit Customer Complaints  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">380837</a> <a href="#">14 Nov 2025</a>	Audit 14 Complaints And Corrective Actions Viamed (30)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
System Generated BSI Audits Calendar BSI Audit Customer Complaints Audit 14 BEFORE starting the Audit you need to update the Processes attached to the Audit. Find the doc in the Document Index Admin Document - Copy and Paste List Processes Per Title replacing them in the current audit. Review Last years Audit see if its still suitable Carry out the Audit, reviewing and commenting on the ISO route map first. Any follow on issues must be identified with Observation Issue or a Non Conformance Issue. If the issue is a Non Conformance a QC21 form must be filled in, as per the QC21 form procedure, and investigated as such. If there are any non Conformances from the Audit, create a follow up / related Issue including all of the following - With a time for Completion Immediate Action Plan Corrective Action Plan Corrective Action Confirmation of Resolution If its a major / critical non conformance complete form QC 18  
H.S.E. implications :No health and safety implications

**25 Nov 2025 Helen Lamb**  
Audit attached, nothing outstanding from the last audit. No viamed issues. There is a VST outstanding follow issue from rolling tasks 381760 task 1068 not up to date. But this is not Viamed No problems no non conformances for Viamed

**26 Nov 2025 Derek Lamb**  
reviewed ok

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">380813</a> <a href="#">13 Nov 2025</a>	Order Error : 160012 Viamed Account error	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb  
Auto Issue from Error Log 160012  
Order Entered by Sophie Lines  
Order Checked by Aqib Majeed  
Office  
Error was Viamed  
Account error  
Fault:  
VAT has been added to order even though it is a thailand order. should not have VAT on it ACC has been set up incorrectly, checker did not spot either, stopped at goods out.

Possible Fix  
passed back to office

**14 Nov 2025 Helen Lamb**  
Done

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		conformance / must state in issue if its an observation						
<a href="#">380348</a> <a href="#">07 Nov 2025</a>	Shipped Items Return to Supplier BOX1054	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
Does this Return BOX1054 warrant a NON conformance report via the CAPA process VM3COP10

**07 Nov 2025 Derek Lamb**  
Leaking gas ? the figure seems unusually high. We're waiting for a response from the supplier.

**24 Feb 2026 Derek Lamb**  
Created Related Issue #389899  
Added by Derek Lamb sent to Derek Lamb  
Original Subject: Shipped Items Return to Supplier BOX1054\_380348.1 Leaking gas ? the figure seems unusually high. We're waiting for a response from the supplier. issue untagged still yellow but low do the list

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">380043</a> <a href="#">05 Nov 2025</a>	Non conformance review history Viamed (283)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
System Generated Check the history of the last Non conformance review, check actions are being carried out, and non conformances are not reoccurring. Review the Order Invoice Error Logs (link in issue) for increases in errors. Areas that are errors that are our mistake and not due to changes instigated by the customer. Note any problem areas in report. Review Customer Complaint and Non Conformance Review Screen check back over the previous month since last review to see if anything internal / staff / process related needs to be highlighted. Go to Add an issue, search non con and review any issues since the last review for internal / staff / process related issues that needs to be highlighted. Note any problems in the issue and send to Derek to review  
H.S.E. implications :No health and safety implications

**07 Nov 2025 Helen Lamb**  
Checked back through all Non Conformance issues (not automatically generated) to 1st November 25. We have a non conformances QC21 form Issues 378839 sensors labelled wrong in goods in. Nothing else of concern or to be investigated or reported on in Non Con issues. Review VIAMED Feedback - Product Feedback Negative (741) nothing new. Review VIAMED Feedback - Product Feedback Positive (1189) nothing new. Review VIAMED Feedback - Customer Feedback Negative (739) nothing new. Review VIAMED Feedback - Customer Feedback Positive (1188) nothing new. Review VIAMED Feedback - Customer Complaints (737) nothing new. Order Invoice Error Logs - only one data entry error no other issue, nothing to worry about. Non Conformance, complaints and feedback headers Positive Feedback 377363 07/10/25 2810049 - feedback via ebay and 378028 14/10/25 Flowsensor A Reprocessing Survey: Medtrest LLC. Nothing else new not already covered. Non Conformance issues review screen - No issues not already reviewed above in log. No other feedback issues relating to non conformances that need to be monitored or reviewed. Derek please review the above

07 Nov 2025 Derek Lamb

thankyou

Audit Analysis of Data  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">380039</a> <a href="#">05 Nov 2025</a>	Review Company Data (114)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb

System Generated Review Sales - Number Invoices / Credits per day reports Search for any potential problems in the Data provided

Employee Menu Check Audit Roles Titles and Processes in Employee Roles and Titles. Check the Page for Red Crosses and potentially missing objectives Search the Outstanding Jobs column for potential problems Do we have enough resources. Check the Issues Statistics in the meeting header, Check Issues are being completed in a timely manner.

H.S.E. implications :No health and safety implications

05 Nov 2025 Derek Lamb

Done

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Issue requiring a corrective action plan					
<a href="#">378839</a> <a href="#">22 Oct 2025</a>	QC 21 Non Conformance Report - We have labelled 25 Maxtec MAX55E 0110452 sensors as MAX250E 01104294, in goods in.	<input type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>

Derek Lamb  
Added by Helen Lamb sent to Derek Lamb  
We have labelled 25 Maxtec MAX55E 0110452 sensors as MAX250E 01104294, in goods in. QC 21 form has been generated and attached. please review and add anything you think i may have missed. Once you have accepted this i'll crack on with the corrective actions  
**22 Oct 2025 Helen Lamb**  
Done  
**22 Oct 2025 Helen Lamb**  
Email being sent out Important: Incorrectly Labelled Oxygen Sensors ? Action Required Inbox Sophie Lines Attachments 14:41 (1 hour ago) to deirdre.evans, cathy.green, Helen.lamb Good afternoon, I'm reaching out regarding your recent order with us ? 70004698. We've identified an issue affecting a batch of oxygen sensors that were incorrectly labelled as MAX-250E, while the sensors inside the packaging are a different type. Unfortunately, the 10x MAX-250E sensors shipped to you on 15th October are part of this batch (please refer to Line 1 on the attached delivery note). To resolve this promptly, we kindly ask that you return the affected units (if still in your possession) at your earliest convenience. A prepaid returns label will be emailed to you shortly for this purpose. We are also arranging for 10x correctly labelled MAX-250E sensors to be sent to you as soon as possible. We sincerely apologise for any inconvenience this may have caused and truly appreciate your cooperation in helping us correct the issue quickly. If you have any questions or need further assistance, please don't hesitate to contact me. Kind regards Sophie Lines Office Administrator Viamed Ltd. Done  
**22 Oct 2025 Helen Lamb**  
Please review  
**22 Oct 2025 Derek Lamb**  
ok  
**24 Oct 2025 Derek Lamb**  
noted  
**07 Nov 2025 Derek Lamb**  
reviewed happy with this,  
**11 Dec 2025 Derek Lamb**  
Done

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		requirement to Escalate Non conformance / must state in issue if its an observation						
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<a href="#">378691</a> <a href="#">21 Oct 2025</a> 378259	Software Validation Non Conformance Product Risk Feedback Loop (789)_378259	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
 Added by Derek Lamb sent to Derek Lamb  
 System Generated Task To be Completed testitem TEST FEEDBACK LOOP Create a Non Conformance from this Issue then review the Post Market Surveillance against the Test Technical file zztest not real file. Scroll down to the Issues section, the Non conformance issue and this Issue Should be present H.S.E. implications :No health and safety implications  
**21 Oct 2025 Derek Lamb**  
 test issue only, works as intened

Audit Organisation and Process Verification Internal Process Verification  
 Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">378348</a> <a href="#">17 Oct 2025</a>	Email Distribution (366)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
 System Generated Check the global mailbox, there should be no mails more than a Day old. Give allowance for Monday mornings, there should be no emails older than Friday afternoon. Also note size of space used in Global inbox  
 H.S.E. implications :No health and safety implications  
**21 Oct 2025 Derek Lamb**  
 emails upto date

Non Conformance Issues  
 Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective	Reviewed Non Conformity / Complaint and determine if its a vigilance	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Issue requiring a corrective action plan					
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<a href="#">378315</a> <a href="#">16 Oct</a> <a href="#">2025</a>	Order Error : 159472 Viamed Checking error	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Helen Lamb  
Auto Issue from Error Log 159472  
Order Entered by Sophie Lines  
Order Checked by Aqib Majeed  
Office  
Error was Viamed  
Checking error  
Fault:  
Uploaded amended order and added the two extra lines of stock, but missed that they had also increased one of the original stock lines. Checker also did not spot this even though the final total did not match

Possible Fix  
new order for the missed product and foc carriage for the error

**21 Oct 2025 Helen Lamb**  
Done

Audit Management Review  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed

<a href="#">378231</a> <a href="#">16 Oct</a> <a href="#">2025</a>	Audit 18 Management Review Viamed (21)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
System Generated Audit 18 BEFORE starting the Audit you need to update the Processes attached to the Audit. Find the doc in the Document Index Admin Document - Copy and Paste List Processes Per Title replacing them in the current audit.

Review Last years Audit see if its still suitable Carry out the Audit, reviewing and commenting on the ISO route map first. Any follow on issues must be identified with Observation Issue or a Non Conformance Issue. If the issue is a Non Conformance a QC21 form must be filled in, as per the QC21 form procedure, and investigated as such. If there are any non Conformances from the Audit, create a follow up / related Issue including all of the following - With a time for Completion Immediate Action Plan Corrective Action Plan Corrective Action Confirmation of Resolution If its a major / critical non conformance complete form QC 18

H.S.E. implications :No health and safety implications

**24 Nov 2025 Helen Lamb**

audit completed nothing outstanding from previous audit. No issues from this audit and no non conformances.

**26 Nov 2025 Derek Lamb**

reviewed ok

Non Conformance Issues

Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">377389</a> <a href="#">07 Oct 2025</a>	Order Error : 159359 Viamed Account error	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb

Auto Issue from Error Log 159359

Order Entered by Aqib Majeed

Order Checked by Zoey Teal

Office

Error was Viamed

Account error

Fault:

wrong prioity

Possible Fix

sent back to office to change

**09 Oct 2025 Helen Lamb**

Done

Non Conformance Issues

Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative	Reviewed Non Conformity / Complaint and determine if its a	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		an observation						
<a href="#">377114</a> <a href="#">06 Oct 2025</a>	Review Company Data (114)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
System Generated Review Sales - Number Invoices / Credits per day reports Search for any potential problems in the Data provided

Employee Menu Check Audit Roles Titles and Processes in Employee Roles and Titles. Check the Page for Red Crosses and potentially missing objectives Search the Outstanding Jobs column for potential problems Do we have enough resources. Check the Issues Statistics in the meeting header, Check Issues are being completed in a timely manner.  
H.S.E. implications :No health and safety implications

**06 Oct 2025 Derek Lamb**  
not many overdue issues: system appear ok, we are having a power outage thursday so need to move the servers this week, maybe tomorrow? Overview :06/10/2025 SectionDate UpdatedValue Viamed Forward Orders02/10/2025£ 63,931.00 Viamed Back Orders02/10/2025£ 193,018.46 Viamed Bank Balance02/10/2025£ 207,618.94 Viamed Debtors02/10/2025£ 209,092.47 Viamed Creditors02/10/2025£ 221,632.82 Barclays Loan to Viamed:02/10/2025£ 0.00 Vandagraph Loan To Viamed:02/10/2025£ 0.00 VST Loan To Viamed:02/10/2025£ 0.00 Viamed Properties Loan to Viamed :£0.00 Loan from GGL JSL to Viamed:02/10/2025£ 0.00 Loan from DL To Viamed:02/10/2025£ 0.00 Loan from SN to Viamed:02/10/2025£ 0.00 VST Bank Balance£ 96843.09 Vandagraph Bank Balance£ 140746.19 Viamed Stock Valuation06/10/2025£ 0 Viamed Turnover01/10/25£ 2,228,216.05 Vst Turnover01/10/25£ 823,758.36 Vand Turnover01/10/25£ 300,009.87 VST Debtors£ 53083 VST Creditors£ 49847 VST Back Orders£ 3689 VST Forward Orders£ 137051 Vandagraph Debtors£ 41785 Vandagraph Creditors£ 20694 Vandagraph Back Orders£ 13331 Vandagraph Forward Orders£ 0 Documents AI Processing StatusTypeCount 0Unprocessed25501 1Processing92 2Completed (no summary)58865 3Failed4690 4Summarizing8 5Summarized3355 6Summary failed344 System Resources FilesystemSizeUsedAvailUse%Mountedon /dev/nvme0n1p21.8T259G1.5T15%/ tmpfs16G016G0%/dev/shm tmpfs3.2G1.8M3.2G1%/run tmpfs5.0M05.0M0%/run/lock /dev/nvme0n1p11.1G6.1M1.1G1%/boot/efi

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">376937</a> <a href="#">02 Oct 2025</a>	Shipped Items Return to Supplier BOX1050	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
Does this Return BOX1050 warrant a NON conformance report via the CAPA process VM3COP10

**03 Oct 2025 Derek Lamb**  
envitec sensors normal fail type no cap required

Supplier Complaints (us to them)  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">376408</a> <a href="#">26 Sep 2025</a>	Mediq purchase order PVM4595 short delivery	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
Added by Steve Nixon sent to Derek Lamb  
23-09-25 Just one box was delivered instead of ten. Also, the goods were not packaged and nor was there a delivery note. There is just a shipping label (no references to P.O. or invoice...) stuck directly on to the box. This is not good as it covers up some of the goods labelling and would be seen as a non conformance by ISO auditors.  
**26 Sep 2025 Steve Nixon**  
Response from Mediq: Hi Steve, Please see below the actions we`ve taken to resolve the issue: The shortage has been reported to our warehouse and transport teams for investigation. Due to the high value of the order, we must complete an investigation. A credit may be issued upon outcome of investigation. If you require the stock urgently, please feel free to place a new order. We apologise for any inconvenience this may have caused you and your team. Kind regards, Carmen Green  
Customer Account Executive  
**09 Jan 2026 Derek Lamb**  
resolved Done

Audit Customer Complaints  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		requirement to Escalate Non conformance / must state in issue if its an observation						
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<a href="#">375505</a> <a href="#">17 Sep 2025</a>	Email Distribution (366)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
System Generated Check the global mailbox, there should be no mails more than a Day old. Give allowance for Monday mornings, there should be no emails older than Friday afternoon. Also note size of space used in Global inbox  
H.S.E. implications :No health and safety implications  
**17 Sep 2025 Derek Lamb**  
main box is upto date

VIAMED Customer Complaints  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">375265</a> <a href="#">12 Sep 2025</a> 375260	Invoice RVM158724-1 8 extra 0110132 supplied by mistake_375260.1	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>
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Catherine Spence  
Added by Derek Lamb sent to Catherine Spence  
Added by Steve Nixon sent to Derek Lamb Ref. invoice RVM158724-1 Ichinen have received 8 x 0110132 R-22AV sensors supplied to them FOC by mistake. They were offered these FOC, but they are unable to accept FOC goods. Serial numbers 2791148, 2791149, 2791150, 2791151, 2791174, 2791175, 2791176, 2791177 Cathy can you scan these barcodes to a new location lable it as Ichinen Complaint  
**12 Sep 2025 Catherine Spence**  
The serial numbers listed in the issue are not the serial numbers but the barcode id numbers , the serial numbers are 277607,277608,277609,277610,277633,277634,277635,277636. I will scan these to a new location now on Location 412497  
**19 Feb 2026 Derek Lamb**  
Was a duplicate Issue, still showing on qc report  
**20 Feb 2026 Catherine Spence**  
noted Done

VIAMED Customer Complaints  
Viamed

Issue / Primary	Subject/Notes	Minor Internal Error i.e. not ISO	Reviewed Non Conformity	Determined Cause of Non	Evaluated action to Ensure	Planning and documenting action needed	Verify Action does not adversely	Effectiveness of corrective
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<a href="#">375175</a> <a href="#">11 Sep</a> <a href="#">2025</a>	BSI Non Conformance 2700663-202509-N2	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Helen Lamb  
 Added by Derek Lamb sent to Helen Lamb  
 The medical devices files process of the organisation is not fully effective, as the organisation has not established requirements to Nonconformity Report Page 4 of 9 ensure that product labelling is up to date and viable. The organisation has not documented labelling requirements as a distributor to verify or maintain product labels implemented and placed by the legal manufacturers are current, compliant and remain legible throughout distribution. Technical files were seen historic, predating the sampled product and existing labels on product was seen not recorded. The product sampled was in relation to Temperature Probe – Skin Contact – Ref 1010132021V – SN 210114478 – Type 0212921 – Viamed Ltd – Produced Oct 2021 – Barcode ID 1856344 – Manufacturer Bluepoint Medical GmbH # Interface Agreement – Bluepoint medical GmbH & Co KG – 21/12/2016  
**11 Sep 2025 Helen Lamb**  
 I have filled in some fields it will need reviewing and adding to but its a start. HL V1

BSI Minor Non conformances

Viamed	Vandagraph	VST	Viamed Properties	The Pointless Logo Company				
Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed

<a href="#">375173</a> <a href="#">11 Sep</a> <a href="#">2025</a>	BSI Non Conformance 2700663-202509-N1	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Helen Lamb  
 Added by Derek Lamb sent to Helen Lamb  
 The internal audit process is not fully effective as raised issues within audit records do not clearly differentiate between observations and non-conformities. Audit records reviewed (Audit 15 and Audit 06) highlighted issues and were raised within the Intrastat (eQMS system). The issues raised did not highlight or state if the issues were an observation or non-conformance. No investigation or CAPA related actions were seen and the issues were cleared/completed with note comment only. # Audit 06 – Performed 30 May 2025 – Issues 365729 Raised 26 June 2025; Issue 367474 Raised 18 June 2025 – Closed 26 June 2025 # Audit 15 – Performed 21 July 2025 – Issues 370409, 370410 – Raised 21 July 2025 – Closed 22 July 2025  
**11 Sep 2025 Helen Lamb**  
 My first version is attached please review it and see if im on the right track. Once we are happy i will start doing the corrective actions

Non Conformance Issues

Viamed								
Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No	Reviewed Non Conformity / Complaint and determine	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed



2025

Derek Lamb  
 System Generated Review Sales - Number Invoices / Credits per day reports Search for any potential problems in the Data provided  
 Employee Menu Check Audit Roles Titles and Processes in Employee Roles and Titles. Check the Page for Red Crosses and potentially missing objectives Search the Outstanding Jobs column for potential problems Do we have enough resources. Check the Issues Statistics in the meeting header, Check Issues are being completed in a timely manner.  
 H.S.E. implications :No health and safety implications  
**11 Sep 2025 Derek Lamb**  
 done

Non Conformance Issues  
 Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">374520</a> <a href="#">04 Sep 2025</a>	Order Error : 158513 Vandagraph New Error	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Helen Lamb  
 Auto Issue from Error Log 158513  
 Order Entered by Aqib Majeed  
 Order Checked by Michael Lamb  
 Goods Out  
 Error was Vandagraph  
 New Error  
 Fault:  
 Order not fulfilled on Shopify  
 Possible Fix  
 Done  
**04 Sep 2025 Helen Lamb**  
 I have chased this again but also added a process and rolling issue to try and stop this happening Done

Audit Documentation Control  
 Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No	Reviewed Non Conformity / Complaint and determine	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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Derek Lamb  
 System Generated Audit 02 Review Last years Audit see if its still suitable Before Proceeding you need to update the Processes attached to the Audit Search the Document in the Index, View the Admin Page Copy and Paste the Attached Processes, replacing them in the current audit Any non Conformances from the Audit: Create a follow up / related Issue, With a time for Completion Immediate Action Plan Corrective Action Plan Corrective Action Confirmation of Resolution if its a major / critical non conformance complete form QC 18  
 H.S.E. implications :No health and safety implications  
**03 Sep 2025 Helen Lamb**  
 Audit completed no non conformances. Nothing outstanding and nothing ongoing. Please review  
**03 Sep 2025 Derek Lamb**  
 thankyou

Audit Purchasing  
 Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">373965</a> <a href="#">01 Sep 2025</a>	Audit 05 Purchasing Suppliers Viamed (37)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
 System Generated Audit 05 BEFORE starting Audit you need to update the Processes attached to the Audit. Find the doc in the Document Index Admin Document - Copy and Paste List Processes Per Title replacing them in the current audit. Review Last years Audit see if its still suitable Carry out the Audit, reviewing and commenting on the ISO route map first. If there are any non Conformances from the Audit, create a follow up / related Issue including all of the following - With a time for Completion Immediate Action Plan Corrective Action Plan Corrective Action Confirmation of Resolution If its a major / critical non conformance complete form QC 18  
 H.S.E. implications :No health and safety implications  
**08 Sep 2025 Helen Lamb**  
 Completed Audit attached, no outstanding issue. No non conformances. Please review  
**09 Sep 2025 Derek Lamb**  
 thankyou

Audit Contract Review  
 Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	corrective action plan					
<a href="#">373964</a> <a href="#">01 Sep 2025</a>	Audit 02 Contract Review Viamed (36)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
System Generated Audit 02 Review Last years Audit see if its still suitable Before Proceeding you need to update the Processes attached to the Audit Search the Document in the Index, View the Admin Page Copy and Paste the Attached Processes, replacing them in the current audit Any non Conformances from the Audit: Create a follow up / related Issue, With a time for Completion Immediate Action Plan Corrective Action Plan Corrective Action Confirmation of Resolution if its a major / critical non conformance complete form QC 18  
H.S.E. implications :No health and safety implications  
**03 Sep 2025 Helen Lamb**  
Audit completed no non conformances. Nothing outstanding and nothing ongoing. Please review  
**03 Sep 2025 Derek Lamb**  
thankyou

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">373734</a> <a href="#">27 Aug 2025</a>	Shipped Items Return to Supplier BOX1044	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
Does this Return BOX1044 warrant a NON conformance report via the CAPA process VM3COP10  
**27 Aug 2025 Derek Lamb**  
8010006 various output errors normal for o2 sensors no cap required recent failure rate (0.31 %)

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">373624</a> <a href="#">27 Aug 2025</a>	Order Error : 158466 Viamed New Error	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Helen Lamb  
Auto Issue from Error Log 158466  
Order Entered by Aqib Majeed  
Order Checked by Sophie Lines  
Non Selected  
Error was Viamed  
New Error  
Fault:  
website error - order not fulfilled on website

Possible Fix  
done

**27 Aug 2025 Helen Lamb**  
Done

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		an observation						
<a href="#">373242</a> <a href="#">20 Aug 2025</a>	Filling in or entering an SRS Please tag them correctly	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb  
 Added by Helen Lamb sent to Helen Lamb  
 INFORMATION ONLY ISSUE **DO NOT ADD NOTES!**  
 Anyone filling in a SRS please can you make sure that you tag them correctly. The repair/quote option is being set to Repair or Service/Calibration, When it should say Quote . It should only be turned from quote to Repair or Service/Calibration when we have the customer purchase order.  
**20 Aug 2025 Helen Lamb**  
 Done

Non Conformance Issues  
 Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">373120</a> <a href="#">19 Aug 2025</a>	Order Error : 158339 Viamed Carriage - office	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Helen Lamb  
 Auto Issue from Error Log 158339  
 Order Entered by Sophie Lines  
 Order Checked by Kate Griffiths  
 Office  
 Error was Viamed  
 Carriage - office  
 Fault:  
 put the export ups express saver on a UK order  
  
 Possible Fix  
 sent back to office for them to amend  
  
**20 Aug 2025 Helen Lamb**  
 done

Non Conformance Issues  
 Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance	Reviewed Non Conformity / Complaint	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and	Verify Action does not adversely affect Safety Performance	Effectiveness of corrective action reviewed
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		or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	and determine if its a vigilance Issue requiring a corrective action plan			implementation QC 28b	or regulatory requirements	
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<a href="#">373092</a> <a href="#">19 Aug 2025</a>	Shipped Items Return to Supplier BOX1043	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
Does this Return BOX1043 warrant a NON conformance report via the CAPA process VM3COP10

**20 Aug 2025 Derek Lamb**  
4410540 The unit has a red ring ISA fault returning to supplier no cap required by us

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">373079</a> <a href="#">19 Aug 2025</a> 363316	Repairs details not being entered correctly	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Helen Lamb  
Added by Robert Connor sent to Helen Lamb  
Still no improvement, it`s the same as ever

**16 Jan 2026 Helen Lamb**  
Derek has added the stock memo to the order entry page and i have also spoken to the office. I hope this gets better Done

**19 Jan 2026 Robert Connor**  
Ok, hopefully that will make a difference, thanks.

Audit Contract Review  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">372880</a> <a href="#">18 Aug 2025</a>	Distributor Agreements (379)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
System Generated  
Note this is an Audit - simply need to ensure its being carried out

Sales -> Distributor Agreements -> Check Sales Against Agreements,

List should be up to date / empty.  
H.S.E. implications :No health and safety implications  
**19 Aug 2025 Derek Lamb**  
some on list but only a few distributors, so the list is being maintained

Audit Organisation and Process Verification Internal Process Verification  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">372877</a> <a href="#">18 Aug</a>	Email Distribution (366)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

2025

Derek Lamb  
System Generated Check the global mailbox, there should be no mails more than a Day old. Give allowance for Monday mornings, there should be no emails older than Friday afternoon. Also note size of space used in Global inbox  
H.S.E. implications :No health and safety implications  
**19 Aug 2025 Derek Lamb**  
main box has been done, all upto date

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">372591</a> <a href="#">13 Aug 2025</a>	Shipped Items Return to Supplier BOX1042	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
Does this Return BOX1042 warrant a NON conformance report via the CAPA process VM3COP10

**13 Aug 2025 Derek Lamb**

Supplier Return ? Summary (BOX?1042) #Stock RefSupply RefBar?codeSerial No.Fault / IssueCountry of OriginCustoms ValueEst. Date inAdded to BOX?1042Added By 10110040??V108401Leaking gasesGermany€?5.0003?Mar?202501?Aug?2025Catherine?Spence 20110040??V108402Leaking gasesGermany€?5.0003?Mar?202501?Aug?2025Catherine?Spence 30110040??V108403Leaking gasesGermany€?5.0003?Mar?202501?Aug?2025Catherine?Spence 40110049??V104860UnstableGermany€?5.0019?Dec?202401?Aug?2025Catherine?Spence 50110182??V118077No OutputGermany€?5.0028?Apr?202501?Aug?2025Catherine?Spence Key Points ItemTotal ValuePrimary IssueNumber of Units 0110040 (3 units)€?15.00Leaking gases3 0110049 (1 unit)€?5.00Unstable1 0110182 (1 unit)€?5.00No output1 Grand Total€?25.00?5 Operations Overview Print RMA Request PDF ? All items are ready for RMA processing; PDFs are queued for printing. Add Items from Print Queue ? The five items above have already been added to the print queue for shipment. Non?Serial/Bar? coded Items ? All units are non?serial and bar?coded, so the system uses ?N/A? for the serial number field where applicable. Customs Information ? Each item is from Germany, with a declared customs value of €?5.00. Shipping Invoice / Fault Sheets ? Fault sheets are generated using the serial numbers provided (e.g., V108401). Boxing / Packaging ? All items are listed under BOX?1042 with the same ?Date Added? (01?Aug?2025) for consistency in the packing list.

**13 Aug 2025 Derek Lamb**

reviewed no cap required

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative	Reviewed Non Conformity / Complaint and determine if its a	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	vigilance Issue requiring a corrective action plan					
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<a href="#">372212</a> <a href="#">08 Aug</a> <a href="#">2025</a>	Shipped Items Return to Supplier BOX1041	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
Does this Return BOX1041 warrant a NON conformance report via the CAPA process VM3COP10

**11 Aug 2025 Derek Lamb**  
sensor output errors, no cap required

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">371760</a> <a href="#">05 Aug</a> <a href="#">2025</a>	Non conformance review history Viamed (283)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
System Generated Check the history of the last Non conformance review, check actions are being carried out, and non conformances are not reoccurring. Review the Order Invoice Error Logs (link in issue) for increases in errors. Areas that are errors that are our mistake and not due to changes instigated by the customer. Note any problem areas in report. Review Customer Complaint and Non Conformance Review Screen check back over the previous month since last review to see if anything internal / staff / process related needs to be highlighted. Go to Add an issue, search non con and review any issues since the last review for internal / staff / process related issues that needs to be highlighted. Note any problems in the issue and send to Derek to review  
H.S.E. implications :No health and safety implications

**08 Aug 2025 Helen Lamb**  
Checked back through all Non Conformance issues (not automatically generated) to 1st August 25. 370242 Manufacturing defect, the sensor is leaking at the O ringno cap required returning to supplier, 370602 Calibrate button not working, Nothing else of concern or to be investigated or reported on in Non Con issues. Review VIAMED Feedback - Product Feedback Negative (741) nothing new. Review VIAMED Feedback - Product Feedback Positive (1189) nothing new. Review VIAMED

Feedback - Customer Feedback Negative (739) nothing new. Review VIAMED Feedback - Customer Feedback Positive (1188) nothing new. Review VIAMED Feedback - Customer Complaints (737) nothing new. Order Invoice Error Logs - No issue, nothing to worry about. Non Conformance, complaints and feedback headers. 368629, 368630, 368631, 368990, 369141, 369316, 369493, 369494, 369991, 370005 errors covered in the error log. Issue 367345 missing sensors from Maxtec. Nothing else and nothing of concerns. Non Conformance issues review screen - No issues not already reviewed above in log.No other feedback issues relating to non conformances that need to be monitored or reviewed.Derek please review the above  
**08 Aug 2025 Derek Lamb**  
 thankyou

Audit Analysis of Data  
 Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">371755</a> 05 Aug 2025	Review Company Data (114)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
 System Generated Review Sales - Number Invoices / Credits per day reports Search for any potential problems in the Data provided

Employee Menu Check Audit Roles Titles and Processes in Employee Roles and Titles. Check the Page for Red Crosses and potentially missing objectives Search the Outstanding Jobs column for potential problems Do we have enough resources. Check the Issues Statistics in the meeting header, Check Issues are being completed in a timely manner.

H.S.E. implications :No health and safety implications

**05 Aug 2025**

done, still need to remove SH from th eprocess logs

**05 Aug 2025 Derek Lamb**

done

Future Reviews - Internal Audits  
 Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested /	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation						
<a href="#">371570</a> <a href="#">01 Aug 2025</a>	Objective Ensure the Audits are performed within a timely manner Review the Tasks and Audits for the Audits Should be no more than 1 outstanding issue for each section	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

General  
01 Aug 2025 Derek Lamb  
audits ongoing

Complaints  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">371545</a> <a href="#">01 Aug 2025</a>	Objective: All complaints to be logged, contained / risk controlled in 10 Days and All complaints to be satisfactorily resolved in 6 months (I)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

General  
01 Aug 2025 Derek Lamb  
no complaints

Audit Repairs and Service  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non	Reviewed Non Conformity /	Determined Cause of Non	Evaluated action to Ensure	Planning and documenting action needed and	Verify Action does not adversely affect Safety	Effectiveness of corrective action reviewed
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		Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Complaint and determine if its a vigilance Issue requiring a corrective action plan	Conformity / Complaint	does not recur	implementation QC 28b	Performance or regulatory requirements	
<a href="#">371427</a> <a href="#">01 Aug 2025</a>	Audit 24 Due Servicing (288)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb

System Generated Audit 24 Due Servicing BEFORE starting the Audit you need to update the Processes attached to the Audit. Find the doc in the Document Index Admin Document - Copy and Paste List Processes Per Title replacing them in the current audit. Review Last years Audit see if its still suitable Carry out the Audit, reviewing and commenting on the ISO route map first. If there are any non Conformances from the Audit, create a follow up / related Issue including all of the following - With a time for Completion Immediate Action Plan Corrective Action Plan Corrective Action Confirmation of Resolution If its a major / critical non conformance complete form QC 18  
H.S.E. implications :No health and safety implications

**15 Aug 2025 Derek Lamb**  
thankyou

Audit Health and Saftey  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">371416</a> <a href="#">01 Aug 2025</a>	Audit 19 Health And Saftey Viamed (13)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb

System Generated Do HSE Audit Audit No 19. Send out HSE Personnel Questionnaire, and the HSE DSE Personnel Questionnaire and reissue message of the day reminding users all HSE Documents are available in Intrastats BEFORE starting the Audit you need to update the Processes attached to the Audit. Find the doc in the Document Index Admin Document - Copy and Paste List Processes Per Title replacing them in the current audit. Review Last years Audit see if its still

suitable Carry out the Audit, reviewing and commenting on the ISO route map first. If there are any non Conformances from the Audit, create a follow up / related Issue including all of the following - With a time for Completion Immediate Action Plan Corrective Action Plan Corrective Action Confirmation of Resolution If its a major / critical non conformance complete form QC 18

H.S.E. implications :No health and safety implications

**29 Aug 2025 Helen Lamb**

Completed Audits attached, No outstanding issues, no non conformances. Please review

**29 Aug 2025 Derek Lamb**

thankyou Done

**Non Conformance Issues**

Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">371116</a> <a href="#">29 Jul 2025</a>	Shipped Items Return to Supplier BOX1039	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb

Does this Return BOX1039 warrant a NON conformance report via the CAPA process VM3COP10

**29 Jul 2025 Derek Lamb**

Here`s a summary of failure reports and related QA activity for the Vandagraph VST Sensors (Part 8010007) covering the period 29 Jul 2023 ? 30 Jul 2025: Product Overview Part Number: 8010007 (VST Oxygen Sensor Module) Total Units Booked In (2 Years): 3,950 Total Units in System: 3,943 Installations span: Multiple SOs with new membrane implementations as of Oct 2024 Main configuration: OOD103 variants with individual sensor serial ranges tracked Fault Summary Global Failures (All Time): Total Global Failures Recorded: 68 faults out of 18,650 units in the system Warranty Failures: 57 (??0.31%) Non-Warranty Failures: 11 (??0.06%) No Fault Found cases: 44 (not included in fault rate) Unique Faults (29 Jul 2023 ? 30 Jul 2025): High Output: 1 Low Output: 2 Unstable Output Signal: 1 All from same product code (8010007), using sensor E1001581 Customer Return Timing (from invoice to failure): Fault Type3 Mo4 Mo10 Mo11 Mo Unstable Output Signal1 High Output1 Low Output11 Batch QA Observations Known batches received between Aug 2023 and May 2025 4 warranty failures found across multiple batches Most returns booked from Feb to May 2024, with no significant cluster after May 2024 Single unit returns tracked with serials, e.g. SN 100016?100027 (SO 263031148) ? Breakdown by Fault Code Fault CodeCountFault TypeNotes No Output11FaultMinor occurrence Low Output6FaultIncludes demo/customer returns Unstable Output Signal34FaultHighest single-code fault Linearity Error2FaultNo recent spike Zero Offset Signal Out Of Tolerance4FaultStable trend Output Cap Detached2FaultNo trend High Output4FaultNo spike PCB Corrosion / Connector Corrosion8No FaultLikely environmental Electrolyte Leakage2FaultUncommon PCB Movement Within Casing2FaultHardware issue Connector - unstable connection1FaultSingle instance Total QA-faulted units (true hardware or performance faults): 68 Non-Conformance Risk Review: Currently not triggered (no fault code exceeds 5% failure rate) Monthly Fault Trends No spikes in any month from Jan to Jun 2025; returns remain low volume and evenly spread. ? Risk Assessment Summary Overall Fault Rate: Very low (<?0.4%) No fault code exceeds 5% threshold ? No non-conformance raised Environmental & assembly issues (corrosion, movement, connector) noted but infrequent Recommendations Continue tracking serial ranges against faults (especially post-Oct 2024 batches with new membranes) Monitor ?Unstable Output Signal? in future returns?has highest fault count globally Document all ?No Fault Found? cases separately to avoid skewing future metrics Audit OOD103-1V and similar variants more closely, as most faults relate to this configuration Let me know if you want this formatted as a report or added into a .docx, .xlsx, or presentation

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">371028</a> <a href="#">28 Jul 2025</a>	Order Error : 158102 Viamed Address Error - Office	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb  
Auto Issue from Error Log 158102  
Order Entered by Ryan Swaine  
Order Checked by Aqib Majeed  
Vandagraph Office  
Error was Viamed  
Address Error - Office  
Fault:  
Postcode/ Zip code missed off order, even though it was in the address listed on the email

Possible Fix

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		an observation						
<a href="#">370655</a> <a href="#">23 Jul</a> <a href="#">2025</a>	Order Error : 157850 Viamed Missing info - office	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb  
Auto Issue from Error Log 157850  
Order Entered by Aqib Majeed  
Order Checked by Emily Hanson  
Office  
Error was Viamed  
Missing info - office  
Fault:  
no PO added and there was one on the paperwork .11072594AM auto generate in intrastats should have been LR746607

Possible Fix  
missed on entry and also on checking

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">370635</a> <a href="#">23 Jul</a> <a href="#">2025</a>	Order Error : 156402 Viamed New Error	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb  
Auto Issue from Error Log 156402  
Order Entered by Aqib Majeed  
Order Checked by Kate Griffiths  
Non Selected  
Error was Viamed  
New Error  
Fault:  
Estimated delivery date not updated as per RocketChat message.

Possible Fix  
If aware delivery date will be earlier, requires to be changed.

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">370634</a> <a href="#">23 Jul</a> <a href="#">2025</a>	Order Error : 157477 Viamed New Error	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Helen Lamb  
Auto Issue from Error Log 157477  
Order Entered by Kate Griffiths  
Order Checked by Aqib Majeed  
Goods Out  
Error was Viamed  
New Error  
Fault:  
Estimated delivery date not updated as per RocketChat message.

Possible Fix  
If aware delivery date will be earlier, requires to be changed.

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		no requirement to Escalate Non conformance / must state in issue if its an observation						
<a href="#">370409</a> <a href="#">21 Jul 2025</a> 362949	Audit 15 Production Viamed (28) Q12 Jobs in Start Job List	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Catherine Spence  
 Added by Helen Lamb sent to Catherine Spence  
 PRO3457 / PS3757 part 0110073 from March is still on the Start Job List in production. please can you review it and see if it should be there

**22 Jul 2025 Catherine Spence**  
 will get derek to clear as we do not make that sensor this way, job not added by me

Non Conformance Issues  
 Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">370384</a> <a href="#">21 Jul 2025</a>	Order Error : 157328 Viamed Missing info - office	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb  
 Auto Issue from Error Log 157328  
 Order Entered by Kate Griffiths  
 Order Checked by Emily Hanson  
 Office  
 Error was Viamed  
 Missing info - office  
 Fault:  
 Customer was told stock 0110805 was due 23/07/25 and was told this on 18/06/25, but the PVM was updated on the 18/06/25 to stated expected Sept 2025, no one had informed customer until today 21/07/25. Customer wanted a part shipment, but due to being proforma , they should pay a second shipment

Possible Fix  
 Agreed to second second carriage cost FOC as we had not informed customer agreed with DL. Also have spoken to office to show them how to stop this from happening again, by watching the PVM on the system, so that any amendments done to the order, it will send an issue to all the people watching that order, and will let them know its been amended, and then they can then contact the customer with an advised lead time if needed.

Supplier Complaints (us to them)  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">370352</a> <a href="#">21 Jul 2025</a>	Maxtec sales order 352188 - 10 x R125P01-007 missing from the shipment	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
Added by Steve Nixon sent to Derek Lamb  
See the attached.

**21 Jul 2025 Steve Nixon**

Supplier Complaints (us to them)  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">370349</a> <a href="#">21 Jul 2025</a>	Maxtec sales order 352188 - 10 x R125P01-007 missing from the shipment	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
 Added by Steve Nixon sent to Derek Lamb  
 See the attached.

**21 Jul 2025 Steve Nixon**  
 Once read/processed this can be closed. 10 sensors were later supplied on 16.06.25

**21 Jul 2025 Derek Lamb**  
 Done

**Non Conformance Issues**  
 Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">370242</a> <a href="#">18 Jul 2025</a>	Shipped Items Return to Supplier BOX1040	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
 Does this Return BOX1040 warrant a NON conformance report via the CAPA process VM3COP10

**21 Jul 2025 Derek Lamb**  
 0110040 Manufacturing defect, the sensor is leaking at the O ring no cap required returning to supplier

**Audit Organisation and Process Verification Internal Process Verification**  
 Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		no requirement to Escalate Non conformance / must state in issue if its an observation						
<a href="#">370029</a> <a href="#">17 Jul</a> <a href="#">2025</a>	Email Distribution (366)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
System Generated Check the global mailbox, there should be no mails more than a Day old. Give allowance for Monday mornings, there should be no emails older than Friday afternoon. Also note size of space used in Global inbox

**18 Jul 2025 Derek Lamb**  
all upto date

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">369991</a> <a href="#">16 Jul</a> <a href="#">2025</a>	Order Error : 157721 Viamed Missing info - office	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Helen Lamb  
Auto Issue from Error Log 157721  
Order Entered by Aqib Majeed  
Order Checked by Kate Griffiths  
Office  
Error was Viamed  
Missing info - office  
Fault:  
No message to state to add their BTL reference to all stock, even though its stated on the Purchase Order  
  
Possible Fix

Audit Health and Saftey  
Viamed

Issue / Primary	Subject/Notes	Minor Internal Error	Reviewed Non	Determined Cause of	Evaluated action to	Planning and documenting	Verify Action does not	Effectiveness of corrective
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		to Escalate Non conformance / must state in issue if its an observation						
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<a href="#">369024</a> <a href="#">07 Jul</a> <a href="#">2025</a>	Non conformance review history Viamed (283)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
System Generated Check the history of the last Non conformance review, check actions are being carried out, and non conformances are not reoccurring. Review the Order Invoice Error Logs (link in issue) for increases in errors. Areas that are errors that are our mistake and not due to changes instigated by the customer. Note any problem areas in report. Review Customer Complaint and Non Conformance Review Screen check back over the previous month since last review to see if anything internal / staff / process related needs to be highlighted. Go to Add an issue, search non con and review any issues since the last review for internal / staff / process related issues that needs to be highlighted. Note any problems in the issue and send to Derek to review

**17 Jul 2025 Helen Lamb**  
Next Action Changed From Helen Lamb To Derek Lamb Checked back through all Non Conformance issues (not automatically generated) to 1st July 25. 368610 R17AH high failure rate, 368629, 368630, 368631. Nothing else of concern or to be investigated or reported on in Non Con issues. Review VIAMED Feedback - Product Feedback Negative (741) nothing new. Review VIAMED Feedback - Product Feedback Positive (1189) nothing new. Review VIAMED Feedback - Customer Feedback Negative (739) nothing new. Review VIAMED Feedback - Customer Feedback Positive (1188) nothing new. Review VIAMED Feedback - Customer Complaints (737) nothing new. Order Invoice Error Logs - No issue, nothing to worry about. Non Conformance, complaints and feedback headers. 368629, 368630, 368631, 368990, 369141, 369316, 369493, 369494, 369991, 370005 errors covered in the error log. Issue 367345 missing sensors from Maxtec. Nothing else and nothing of concerns. Non Conformance issues review screen - No issues not already reviewed above in log. No other feedback issues relating to non conformances that need to be monitored or reviewed. Derek please review the above

**18 Jul 2025 Derek Lamb**  
Done

Audit Analysis of Data  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">369017</a> <a href="#">07 Jul</a> <a href="#">2025</a>	Review Company Data (114)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
System Generated Review Sales - Number Invoices / Credits per day reports Search for any potential problems in the Data provided

Employee Menu Check Audit Roles Titles and Processes in Employee Roles and Titles. Check the Page for Red Crosses and potentially missing objectives Search the Outstanding Jobs column for potential problems Do we have enough resources. Check the Issues Statistics in the meeting header, Check Issues are being completed in a timely manner.

**07 Jul 2025 Derek Lamb**

Done

Non Conformance Issues

Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">368990</a> <a href="#">04 Jul 2025</a>	Order Error : 157677 Vandagraph Checking error	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb

Auto Issue from Error Log 157677

Order Entered by Ryan Swaine

Order Checked by Kate Griffiths

Vandagraph Office

Error was Vandagraph

Checking error

Fault:

Down as a pri 3 but with recorded delivery line for carriage , if its a true recorded delivery then it shouyld be a pri 4

Possible Fix

passed back to office to correct

Non Conformance Issues

Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	corrective action plan						
<a href="#">368629</a> <a href="#">02 Jul 2025</a>	0110137 QA Data Requires Risk / Non Conformance Review Jun 2025	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
Added by Robert Connor sent to Derek Lamb  
INFORMATION ONLY ISSUE **DO NOT** ADD NOTES!  
QA Failures High Numbers

**Audit Repairs and Service Viamed**

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">368451</a> <a href="#">01 Jul 2025</a>	Audit 11 Repairs And Service Viamed (171)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
System Generated Audit 11 BEFORE starting the Audit you need to update the Processes attached to the Audit. Find the doc in the Document Index Admin Document - Copy and Paste List Processes Per Title replacing them in the current audit. Review Last years Audit see if its still suitable Carry out the Audit, reviewing and commenting on the ISO route map first. If there are any non Conformances from the Audit, create a follow up / related Issue including all of the following - With a time for Completion Immediate Action Plan Corrective Action Plan Corrective Action Confirmation of Resolution If its a major / critical non conformance complete form QC 18

**24 Jul 2025 Helen Lamb**  
Next Action Changed From Helen Lamb To Derek Lamb completed audit attached no outstanding issues no non conformances. Nothing outstanding from last audit. Please review

**24 Jul 2025 Derek Lamb**  
Done

Audit Training Viamed								
Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">368315</a> <a href="#">30 Jun</a> <a href="#">2025</a>	Audit 08 Training Viamed (10)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
System Generated Audit 08 Training BEFORE starting the Audit you need to update the Processes attached to the Audit. Find the doc in the Document Index Admin Document - Copy and Paste List Processes Per Title replacing them in the current audit. Review Last years Audit see if its still suitable Carry out the Audit, reviewing and commenting on the ISO route map first. If there are any non Conformances from the Audit, create a follow up / related Issue including all of the following - With a time for Completion Immediate Action Plan Corrective Action Plan Corrective Action Confirmation of Resolution If its a major / critical non conformance complete form QC 18

**11 Jul 2025 Derek Lamb**  
Next Action Changed From Derek Lamb To Helen Lamb please update the check list print and return to me to fill in thankyou

**22 Jul 2025 Helen Lamb**  
Done and printed for you

**24 Jul 2025 Helen Lamb**  
Next Action Changed From Helen Lamb To Derek Lamb ive scanned the audit as requested. Please find it attached, i do not see any outstanding issues.

**24 Jul 2025 Derek Lamb**  
Done

Non Conformance Issues Viamed								
Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested /	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed

		Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation						
<a href="#">368312</a> <a href="#">27 Jun 2025</a>	UnShipped Items Return to Supplier BOX1026	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
Does this Return BOX1026 warrant a NON conformance report via the CAPA process VM3COP10

**03 Jul 2025 Derek Lamb**  
Maxtec Oxygen sensor MAX-250E not returned, no cap required

Audit Calibration  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">367474</a> <a href="#">18 Jun 2025</a> 364782	Audit 06 Calibration Viamed (20) Need to make sure labels Indication Only NOT Calibrated	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Philip Crossley  
Added by Helen Lamb sent to Philip Crossley  
Need to make sure labels Indication Only NOT Calibrated are available. These are for the items in the calibration index that are not calibrated. More labels coming tomorrow so they can continue to be used.

**19 Jun 2025 Philip Crossley**  
The stickers have arrived so I have stickered the devices from Lindas area and the DVM. I will have a look around next week to see if I have missed anything.

**25 Jun 2025 Philip Crossley**  
I have stickered a few more things in Michael Greens room. I think Im probably done, if you come across any more I have plenty of stickers left.

Complaints from Viamed to a Supplier  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">367346</a> <a href="#">17 Jun 2025</a>	Maxec shpment - 10 missing MAX-250 oxygen sensors	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
 Added by Steve Nixon sent to Derek Lamb  
 From shipment dated 02-06-2025 10 of the R125P01-007 MAX-250 sensors are missing (serial numbers LD30499051 to LD30499060) We have received 40 sensors instead of 50 See the attached documents.

**17 Jun 2025 Derek Lamb**

**03 Jul 2025 Derek Lamb**  
noted

Complaints from Viamed to a Supplier  
 Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">367345</a> <a href="#">17 Jun 2025</a>	Maxec shpment - 10 missing MAX-250 oxygen sensors	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
 Added by Steve Nixon sent to Derek Lamb

From shipment dated 02-06-2025 10 of the R125P01-007 MAX-250 sensors are missing (serial numbers LD30499051 to LD30499060) We have received 40 sensors instead of 50 See the attached documents.

**17 Jun 2025 Derek Lamb**  
duplicate issue

Audit Organisation and Process Verification Internal Process Verification  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">367248</a> <a href="#">17 Jun 2025</a>	Email Distribution (366)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
System Generated Check the global mailbox, there should be no mails more than a Day old. Give allowance for Monday mornings, there should be no emails older than Friday afternoon. Also note size of space used in Global inbox

**17 Jun 2025 Derek Lamb**  
emails inbox all upto date

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		an observation						
<a href="#">366954</a> <a href="#">12 Jun 2025</a>	Order Error : 157293 Viamed Missing info - office	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb  
Auto Issue from Error Log 157293  
Order Entered by Kate Griffiths  
Order Checked by Emily Hanson  
Office  
Error was Viamed  
Missing info - office  
Fault:  
Missed contact number off order, and checker did not spot it either.

Possible Fix

**Audit Documentation Control**  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">366637</a> <a href="#">10 Jun 2025</a>	Audit 10 Documentation Control Viamed (27)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
System Generated BSI Audits Calendar BSI Audit Documentation Control Audit 10. BEFORE starting the Audit you need to update the Processes attached to the Audit. Find the doc in the Document Index Admin Document - Copy and Paste List Processes Per Title replacing them in the current audit. Review Last years Audit see if its still suitable Carry out the Audit, reviewing and commenting on the ISO route map first. If there are any non Conformances from the Audit, create a follow up / related Issue including all of the following - With a time for Completion Immediate Action Plan Corrective Action Plan Corrective Action Confirmation of Resolution If its a major / critical non conformance complete form QC 18

**13 Jun 2025 Helen Lamb**  
Next Action Changed From Helen Lamb To Derek Lamb completed audit attached no ongoing issues. No issues outstanding from last time and no non conformances. Please review

**16 Jun 2025 Derek Lamb**  
thankyu

**Non Conformance Issues**  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">366597</a> <a href="#">09 Jun</a> <a href="#">2025</a>	Order Error : 157193 Viamed Checking error	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Helen Lamb  
Auto Issue from Error Log 157193  
Order Entered by Aqib Majeed  
Order Checked by Kate Griffiths  
Office  
Error was Viamed  
Checking error  
Fault:  
Should be on a pri8 as we are unable to send Teledyne sensors to uk customers for the time being

Possible Fix  
passed back to office

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		/ must state in issue if its an observation						
<a href="#">366217</a> <a href="#">05 Jun 2025</a>	Non conformance review history Viamed (283)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
System Generated Check the history of the last Non conformance review, check actions are being carried out, and non conformances are not reoccurring. Review the Order Invoice Error Logs (link in issue) for increases in errors. Areas that are errors that are our mistake and not due to changes instigated by the customer. Note any problem areas in report. Review Customer Complaint and Non Conformance Review Screen check back over the previous month since last review to see if anything internal / staff / process related needs to be highlighted. Go to Add an issue, search non con and review any issues since the last review for internal / staff / process related issues that needs to be highlighted. Note any problems in the issue and send to Derek to review

**05 Jun 2025 Helen Lamb**  
Next Action Changed From Helen Lamb To Derek Lamb Checked back through all Non Conformance issues (not automatically generated) to 1st June 25. 363312, 363313, 363314, 363315, 365824, 365873, 365874, 365875 high failure from Goods in. Normal issues. Nothing else of concern or to be investigated or reported on in Non Con issues. Review VIAMED Feedback - Product Feedback Negative (741) nothing new. Review VIAMED Feedback - Product Feedback Positive (1189) nothing new. Review VIAMED Feedback - Customer Feedback Negative (739) nothing new. Review VIAMED Feedback - Customer Feedback Positive (1188) nothing new. Review VIAMED Feedback - Customer Complaints (737) nothing new. Order Invoice Error Logs - No issue, nothing to worry about. Non Conformance, complaints and feedback headers 366269 positive feedback. 366297, 366296, 366209, 366206 returns boxes. 365875, 365874, 365873 review of stock. Nothing else and nothing of concerns. Non Conformance issues review screen - No issues not already reviewed above in log. No other feedback issues relating to non conformances that need to be monitored or reviewed. Derek please review the above

**06 Jun 2025 Derek Lamb**  
thankyou

**Audit Analysis of Data Viamed**

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">366211</a> <a href="#">05 Jun 2025</a>	Review Company Data (114)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
System Generated Review Sales - Number Invoices / Credits per day reports Search for any potential problems in the Data provided

Employee Menu Check Audit Roles Titles and Processes in Employee Roles and Titles. Check the Page for Red Crosses and potentially missing objectives Search the Outstanding Jobs column for potential problems Do we have enough resources. Check the Issues Statistics in the meeting header, Check Issues are being completed in a timely manner.

**11 Jun 2025 Derek Lamb**  
 reviewed, re allocated steve hardaker issues

Non Conformance Issues  
 Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">366209</a> <a href="#">04 Jun 2025</a>	UnShipped Items Return to Supplier BOX1022	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
 Does this Return BOX1022 warrant a NON conformance report via the CAPA process VM3COP10

**05 Jun 2025 Derek Lamb**  
 automotive sensors failed qa, going back to supplier, no cap required

Non Conformance Issues  
 Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">366206</a> <a href="#">04 Jun</a>	UnShipped Items Return to Supplier	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

2025 | BOX1028

Derek Lamb  
Does this Return BOX1028 warrant a NON conformance report via the CAPA process VM3COP10

**05 Jun 2025 Derek Lamb**  
automotive sensors failed qa, going back to supplier, no cap required

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">365875</a> <a href="#">02 Jun 2025</a>	0110142 QA Data Requires Risk / Non Conformance Review May 2025	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
Added by Robert Connor sent to Derek Lamb  
INFORMATION ONLY ISSUE **DO NOT** ADD NOTES!  
QA Failures High Numbers

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		Non conformance / must state in issue if its an observation						
<a href="#">365729</a> <a href="#">30 May 2025</a> 364782	Audit 06 Calibration Viamed (20) Checking areas for calibration equipment that does not have a sticker	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Michael Green  
 Added by Helen Lamb sent to Michael Green  
 Checking areas for calibration equipment that does not have a sticker. Please review your areas and look for calibration equipment that does not have a identifying calibration label. Any found unidentified need to be added please.

**11 Jun 2025 Michael Green**  
 I believe all my items are up to date.

Non Conformance Issues  
 Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">364967</a> <a href="#">21 May 2025</a>	Order Error : 156792 Viamed Carriage - goods out	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Helen Lamb  
 Auto Issue from Error Log 156792  
 Order Entered by Kate Griffiths  
 Order Checked by Aqib Majeed  
 Goods Out  
 Error was Viamed  
 Carriage - goods out  
 Fault:  
 Missed two serial numbers off customer paperwork

Possible Fix  
 Given to helen to ammend paperwork

Non Conformance Issues  
 Viamed

Issue / Primary	Subject/Notes	Minor Internal Error	Reviewed Non	Determined Cause of	Evaluated action to	Planning and documenting	Verify Action does not	Effectiveness of corrective
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ID / Call ID		i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Non Conformity / Complaint	Ensure does not recur	action needed and implementation QC 28b	adversely affect Safety Performance or regulatory requirements	action reviewed
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<a href="#">364941</a> <a href="#">21 May 2025</a>	Shipped Items Return to Supplier BOX1032	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
Does this Return BOX1032 warrant a NON conformance report via the CAPA process VM3COP10

**21 May 2025 Derek Lamb**  
oxygen sensor normal fail types, no cap required

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">364934</a> <a href="#">21 May 2025</a>	Shipped Items Return to Supplier BOX1030	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
Does this Return BOX1030 warrant a NON conformance report via the CAPA process VM3COP10

**21 May 2025 Derek Lamb**  
oxygen sensors , normal faults no cap required

Audit Calibration  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">364782</a> <a href="#">20 May 2025</a>	Audit 06 Calibration Viamed (20)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb

System Generated Please Complete Calibration Audit 6 BEFORE starting Audit you need to update the Processes attached to the Audit. Find the doc in the Document Index Admin Document - Copy and Paste List Processes Per Title replacing them in the current audit. Review Last years Audit see if its still suitable Carry out the Audit, reviewing and commenting on the ISO route map first. If there are any non Conformances from the Audit, create a follow up / related Issue including all of the following - With a time for Completion Immediate Action Plan Corrective Action Plan Confirmation of Resolution If its a major / critical non conformance complete form QC 18

**30 May 2025 Helen Lamb**

Created Related Issue #365729

Added by Helen Lamb sent to Michael Green

Checking areas for calibration equipment that does not have a sticker. Please review your areas and look for calibration equipment that does not have a identifying calibration label. Any found unidentified need to be added please.

**18 Jun 2025 Helen Lamb**

Created Related Issue #367474

Added by Helen Lamb sent to Philip Crossley

Need to make sure labels Indication Only NOT Calibrated are available. These are for the items in the calibration index that are not calibrated. More labels coming tomorrow so they can continue to be used.

**19 Jun 2025 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb Audit completed issue sent 367474 to PC and HL to order more Indication only not calibrated stickers to put on equipment that is not calibrated and therefore does not have a date. We have run out. No other issues. nothing outstanding from the last audit and no non conformances.

**19 Jun 2025 Derek Lamb**

thankyou

Audit Contract Review  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective	Reviewed Non Conformity / Complaint and determine if its a vigilance	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Issue requiring a corrective action plan					
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<a href="#">364602</a> <a href="#">19 May 2025</a>	Distributor Agreements (379)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
System Generated  
Note this is an Audit - simply need to ensure its being carried out

Sales -> Distributor Agreements -> Check Sales Against Agreements,  
List should be up to date / empty.

**19 May 2025 Derek Lamb**  
List all Allocated Distributors PulmocorCID10477 PO Portugal 01 Jan 1970 0110701 list is being maintained

Audit Organisation and Process Verification Internal Process Verification  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">364599</a> <a href="#">19 May 2025</a>	Email Distribution (366)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
System Generated Check the global mailbox, there should be no mails more than a Day old. Give allowance for Monday mornings, there should be no emails older than Friday afternoon. Also note size of space used in Global inbox

**19 May 2025 Derek Lamb**  
emails upto date

Non Conformance Issues

Viamed

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<a href="#">364197</a> <a href="#">13 May 2025</a> 339868	Quantity discounts not applied to a quote	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
Added by Derek Lamb sent to Derek Lamb

**03 Jul 2025 Derek Lamb**  
Done

Non Conformance Issues  
Viamed

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<a href="#">363890</a> <a href="#">09 May 2025</a>	UnShipped Items Return to Supplier BOX1014	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
Does this Return BOX1014 warrant a NON conformance report via the CAPA process VM3COP10

13 May 2025 Derek Lamb

jiko and automotive sensors no cap required

Audits Closure

Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">363574</a> <a href="#">07 May 2025</a>	Objective To agree the current schedule of Top Level audits.	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

General

07 May 2025 Derek Lamb

All agree to current schedule of top level audits.

All agreed with 2024 audits results. Closed 2024 audits.

Results of internal audits / Mini Audits

Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">363572</a> <a href="#">07 May</a>	Objective Review the results of internal	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

2025	audits / Mini Audits over the last 12 months. Check everyone is happy with the status of the current years Audits. Reviewed Audit scheduled for next year.							
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General  
07 May 2025 Derek Lamb  
Reviewed mini Audits all happy.  
Reviewed internal Audits all happy.  
Reviewed Mini Audit Management Reviews Global.  
Reviewed Mini Audit Reviews outstanding Issues.  
Reviewed Main Audits 2024.  
Reviewed VST Audit Calendar 2025.  
Discussed follow up issues to 2024 Main Audits.

ISO 9001:2015+A1:2024(E) is Current as per our certification.  
All agreed happy with current years Audits.  
Reviewed all Audits carried out.  
All agreed to continue with existing Audit calendar and scheduled for 2025.

Next BSI Audit will be March 2025.

We reviewed all reports live on the system, nothing flagged up in 2024 audits.  
Follow on Issues

Non Conformities Review  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">363540</a> <a href="#">07 May 2025</a>	Objective To review and action on any non conformances within 15 days. Check for new qc 21 forms.	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

General  
07 May 2025 Derek Lamb  
Nothing new in the Non conformance stock transfer, Non conformance Audit, Non conformance review headings.  
Returns boxes are now showing they are not non conformance issues.  
No outstanding non conformance.

There are no internal QC21 forms.

No issues, all happy with current.

Non Conformance Issues  
Viamed

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<a href="#">363482</a> <a href="#">07 May 2025</a> 354650	Discontinue/Supersede Stock Once Depleted (971) Non Conformance	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
Added by Catrin Hird sent to Helen Lamb  
Issue not been dealt with for at least 3 months.

**09 May 2025 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb can you review and investigate this please

**22 May 2025 Catrin Hird** Resolved

VIAMED Customer Complaints  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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Robert Connor  
 Added by Robert Connor sent to Helen Lamb  
 INFORMATION ONLY ISSUE **DO NOT** ADD NOTES!  
 There have been multiple repairs in the last few months where the repair/quote option has been incorrectly set to Repair or Service/Calibration without any justification. I am repeatedly having to change SRS sheets to Quote when I receive them, as evidence of any customer purchase orders have not been entered correctly into the system. The most recent example of this is SRS69084 Leicester Royal Infirmary, notes indicate a purchase order has been sent but the SRS system has not been updated correctly, so I have changed the repair to "Quote before repair". When I have to do this it delays repairs as the devices will not be serviced/repaired until a purchase order is properly confirmed.  
**13 Aug 2025 Helen Lamb**  
 did this get any better

**19 Aug 2025 Robert Connor**  
 Created Related Issue #373079  
 Added by Robert Connor sent to Helen Lamb  
 Still no improvement, it`s the same as ever

Non Conformance Issues  
 Viamed

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<a href="#">363315</a> <a href="#">06 May 2025</a>	0110137 QA Data Requires Risk / Non Conformance Review Apr 2025	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
 Added by Robert Connor sent to Derek Lamb  
 INFORMATION ONLY ISSUE **DO NOT** ADD NOTES!  
 QA Failures High Numbers

Non Conformance Issues  
 Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	action plan					
<a href="#">363314</a> <a href="#">06 May 2025</a>	0110132 QA Data Requires Risk / Non Conformance Review Apr 2025	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
Added by Robert Connor sent to Derek Lamb  
INFORMATION ONLY ISSUE **DO NOT** ADD NOTES!  
QA Failures High Numbers

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">363313</a> <a href="#">06 May 2025</a>	0110121 QA Data Requires Risk / Non Conformance Review Apr 2025	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
Added by Robert Connor sent to Derek Lamb  
INFORMATION ONLY ISSUE **DO NOT** ADD NOTES!  
QA Failures High Numbers

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective	Reviewed Non Conformity / Complaint and determine if its a vigilance	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Issue requiring a corrective action plan					
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<a href="#">363312</a> <a href="#">06 May 2025</a>	0110107 QA Data Requires Risk / Non Conformance Review Apr 2025	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
 Added by Robert Connor sent to Derek Lamb  
 INFORMATION ONLY ISSUE **DO NOT** ADD NOTES!  
 QA Failures High Numbers

Non Conformance Issues  
 Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">363194</a> <a href="#">05 May 2025</a>	Non conformance review history VST (285)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
 System Generated Check the history of the last Non conformance review, check actions are being carried out, and non conformances are not reoccurring. Review the Order Invoice Error Logs (link in issue) for increases in errors. Areas that are errors that are our mistake and not due to changes instigated by the customer. Note any problem areas in report. Review Customer Complaint and Non Conformance Review Screen check back over the previous month since last review to see if anything internal / staff / process related needs to be highlighted. Go to Add an issue, search non con and review any issues since the last review for internal / staff / process related issues that needs to be highlighted. Note any problems in the issue and send to Derek to review

**19 May 2025 Helen Lamb**  
 Next Action Changed From Helen Lamb To Derek Lamb Checked back through all Non Conformance issues (not automatically generated) to 1st May 25. Of concern or to be investigated or reported on in Non Con issues. Review VST Product Feedback Negative (742) nothing new. Review VST Product Feedback Positive (1190) nothing new. Review VST Feedback - Customer Feedback Negative (740) nothing new. Review VST Customer Feedback Positive(1191) nothing new.

Review VST Feedback - Customer Complaints (738) nothing new. Order Invoice Error Logs - only one data entry error no other issue, nothing to worry about. Non Conformance, complaints and feedback headers nothing new no concerns. Non Conformance issues review screen - No issues not already reviewed above in log. No other feedback issues relating to non conformances that need to be monitored or reviewed. Derek please review the above

**19 May 2025 Derek Lamb**  
thankyou

**Audit Analysis of Data**  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">363187</a> <a href="#">05 May 2025</a>	Review Company Data (114)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
System Generated Review Sales - Number Invoices / Credits per day reports Search for any potential problems in the Data provided

Employee Menu Check Audit Roles Titles and Processes in Employee Roles and Titles. Check the Page for Red Crosses and potentially missing objectives Search the Outstanding Jobs column for potential problems Do we have enough resources. Check the Issues Statistics in the meeting header, Check Issues are being completed in a timely manner.

**07 May 2025 Derek Lamb**  
reviewed

**Non Conformance Issues**  
Viamed

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		to Escalate Non conformance / must state in issue if its an observation						
<a href="#">363027</a> <a href="#">01 May 2025</a>	Order Error : 156056 Vandagraph New Error	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb  
Auto Issue from Error Log 156056  
Order Entered by Kate Griffiths  
Order Checked by Emily Hanson  
Goods Out  
Error was Vandagraph  
New Error  
Fault:  
Order not fulfilled on website

Possible Fix  
Done

Audit Production  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">362949</a> <a href="#">01 May 2025</a>	Audit 15 Production Viamed (28)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
System Generated Audit 15 Production BEFORE starting the Audit you need to update the Processes attached to the Audit. Find the doc in the Document Index Admin Document - Copy and Paste List Processes Per Title replacing them in the current audit. Review Last years Audit see if its still suitable Carry out the Audit, reviewing and commenting on the ISO route map first. If there are any non Conformances from the Audit, create a follow up / related Issue including all of the following - With a time for Completion Immediate Action Plan Corrective Action Plan Corrective Action Confirmation of Resolution If its a major / critical non conformance complete form QC 18

**21 Jul 2025 Helen Lamb**  
Created Related Issue #370409  
Added by Helen Lamb sent to Catherine Spence  
PRO3457 / PS3757 part 0110073 from March is still on the Start Job List in production. please can you review it and see if it should be there

**21 Jul 2025 Helen Lamb**

Created Related Issue #370410  
 Added by Helen Lamb sent to Derek Lamb  
 Production in Production List, in production. Has PRO3719 / PO3718 sent for EMC testing Can you review this and see if it should still being the list and why its there

**22 Jul 2025 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb competed audit attached, no issues from the last audit no non conformance issues. Two issues outstanding, neither are none conformances. Please review

**23 Jul 2025 Derek Lamb**

thankyou

**Audit Post Marketing Surveillance**  
 Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">362859</a> <a href="#">30 Apr 2025</a>	Audit 22 Post Market Surveillance Viamed (14)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

**Helen Lamb**

System Generated Audit 22 BSI Audits Calendar BSI Audit Post Marketing Surveillance BEFORE starting the Audit you need to update the Processes attached to the Audit. Find the doc in the Document Index Admin Document - Copy and Paste List Processes Per Title replacing them in the current audit. Review Last years Audit see if its still suitable Carry out the Audit, reviewing and commenting on the ISO route map first. If there are any non Conformances from the Audit, create a follow up / related Issue including all of the following - With a time for Completion Immediate Action Plan Corrective Action Plan Corrective Action Confirmation of Resolution If its a major / critical non conformance complete form QC 18

**23 May 2025 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb Issue 365215 sent as some rolling issue are out of terms. No other problems nothing outstanding from previous audit and no non conformances. Please find completed audit attached.

**27 May 2025 Derek Lamb**

thnks Done

**27 May 2025 Derek Lamb**

wrong file vst / viamed

**27 May 2025 Derek Lamb**

Next Action Changed From Derek Lamb To Helen Lamb

**27 May 2025 Derek Lamb**

Done

Non Conformance Issues Viamed								
Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">362036</a> <a href="#">19 Apr 2025</a>	Shipped Items Return to Supplier BOX914	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
Does this Return BOX914 warrant a NON conformance report via the CAPA process VM3COP10

**23 Apr 2025 Derek Lamb**  
o2 sensors being return no cap required

Non Conformance Issues Viamed								
Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">362035</a> <a href="#">19 Apr 2025</a>	Shipped Items Return to Supplier BOX1013	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
Does this Return BOX1013 warrant a NON conformance report via the CAPA process VM3COP10

23 Apr 2025 Derek Lamb

o2 sensors being returned , no cap required

Non Conformance Issues

Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">362034</a> <a href="#">19 Apr 2025</a>	Shipped Items Return to Supplier BOX913	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb

Does this Return BOX913 warrant a NON conformance report via the CAPA process VM3COP10

23 Apr 2025 Derek Lamb

o2 sensors, normal sensor fail types

Audit Organisation and Process Verification Internal Process Verification

Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">361858</a> <a href="#">17 Apr</a>	Email Distribution (366)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

2025

Derek Lamb  
System Generated Check the global mailbox, there should be no mails more than a Day old. Give allowance for Monday mornings, there should be no emails older than Friday afternoon. Also note size of space used in Global inbox

17 Apr 2025 Derek Lamb  
working, all upto date

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">361841</a> <a href="#">16 Apr 2025</a>	Order Error : 156300 Vandagraph Carriage - office	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb  
Auto Issue from Error Log 156300  
Order Entered by Aqib Majeed  
Order Checked by Emily Hanson  
Office  
Error was Vandagraph  
Carriage - office  
Fault:  
put at PPRD, but its an export order and should be PPRMITS  
  
Possible Fix  
Sent back to office

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	action plan						
<a href="#">361763</a> <a href="#">16 Apr 2025</a>	Product training not for purpose	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
Added by Robert Connor sent to Helen Lamb  
Not sure what category this belongs under so I'm putting this here. The current method of having Catrin use an LLM to generate quizzes to test training competency is not fit for purpose for two reasons. Catrin has no better training on this than most of us, and LLMs routinely make stuff up because they don't actually understand language. The current quiz for Radiant Warmers is the third round of training in which I have had to inform Catrin that the quiz contains a question that is not answerable, in this case because it has 3 valid answers. Previous quizzes have had questions with no valid answers, and questions where the LLM made up a fact for a question which made it impossible to answer. The way training and testing is done to be reviewed, to ensure that it is done by someone who understands the stock lines in question.

**23 May 2025 Helen Lamb**  
Next Action Changed From Helen Lamb To Derek Lamb

**27 May 2025 Derek Lamb**  
I will discuss this with catrin ,

**27 May 2025 Robert Connor**  
Noted, thanks

**Audit Handling and Storage**  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">361544</a> <a href="#">15 Apr 2025</a>	Audit 07 Handling And Storage Viamed (25)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
System Generated Audit 07 handling and stock control BEFORE starting the Audit you need to update the Processes attached to the Audit. Find the doc in the Document Index Admin Document - Copy and Paste List Processes Per Title replacing them in the current audit. Review Last years Audit see if its still suitable Carry out the Audit, reviewing and commenting on the ISO route map first. If there are any non Conformances from the Audit, create a follow up / related Issue

including all of the following - With a time for Completion Immediate Action Plan Corrective Action Plan Corrective Action Confirmation of Resolution If its a major / critical non conformance complete form QC 18

**23 May 2025 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb completed audit attached no on going issues, non issues from the last audit and no non conformances Please review

**27 May 2025 Derek Lamb**

thankyou Done

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">361144</a> <a href="#">09 Apr 2025</a>	Shipped Items Return to Supplier BOX902	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb

Does this Return BOX902 warrant a NON conformance report via the CAPA process VM3COP10

**10 Apr 2025 Derek Lamb**

automotive sensors failed in QA going back to supplier Done

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		no requirement to Escalate Non conformance / must state in issue if its an observation						
<a href="#">360739</a> <a href="#">07 Apr 2025</a>	Non conformance review history Viamed (283)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb  
System Generated Check the history of the last Non conformance review, check actions are being carried out, and non conformances are not reoccurring. Review the Order Invoice Error Logs (link in issue) for increases in errors. Areas that are errors that are our mistake and not due to changes instigated by the customer. Note any problem areas in report. Review Customer Complaint and Non Conformance Review Screen check back over the previous month since last review to see if anything internal / staff / process related needs to be highlighted. Go to Add an issue, search non con and review any issues since the last review for internal / staff / process related issues that needs to be highlighted. Note any problems in the issue and send to Derek to review

**08 Apr 2025 Helen Lamb**  
Checked back through all Non Conformance issues (not automatically generated) to 1st April 25. Issue 360064 account issue from customer. Resolve nothing ongoing, Issue360344 (360357) high failure rate 0110107. Nothing else of concern or to be investigated or reported on in Non Con issues. Review VIAMED Feedback - Product Feedback Negative (741) nothing new. Review VIAMED Feedback - Product Feedback Positive (1189) nothing new. Review VIAMED Feedback - Customer Feedback Negative (739) nothing new. Review VIAMED Feedback - Customer Feedback Positive (1188) nothing new. Review VIAMED Feedback - Customer Complaints (737) nothing new. Order Invoice Error Logs - No issue, nothing to worry about. Non Conformance, complaints and feedback headers nothing else and nothing of concerns. Non Conformance issues review screen - No issues not already reviewed above in log. No other feedback issues relating to non conformances that need to be monitored or reviewed. Derek please review the above

**Audit Analysis of Data Viamed**

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">360733</a> <a href="#">07 Apr 2025</a>	Review Company Data (114)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
System Generated Review Sales - Number Invoices / Credits per day reports Search for any potential problems in the Data provided  
Employee Menu Check Audit Roles Titles and Processes in Employee Roles and Titles. Check the Page for Red Crosses and potentially missing objectives Search the Outstanding Jobs column for potential problems Do we have enough resources. Check the Issues Statistics in the meeting header, Check Issues are being completed in a timely manner.



2025	Conformance Review Mar 2025							
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Derek Lamb  
 Added by Robert Connor sent to Derek Lamb  
 INFORMATION ONLY ISSUE **DO NOT** ADD NOTES!  
 QA Failures High Numbers

Non Conformance Issues  
 Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">360064</a> <a href="#">31 Mar 2025</a> 360019	Customer Complaints (728) Non Conformance	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
 Added by Derek Lamb sent to Derek Lamb  
 call log ID 5655, Customer has called in response to an email he has received from Zoey saying he has an invoice overdue for £518.40 - he says that is not correct, and he should actually be in credit. Helen to look into when back from annual leave tomorrow.email back and sorted.

**20 May 2025 Derek Lamb**  
 was not really a complaint, was dealt with , helen fixed when returned from holiday

Audit Customer Complaints  
 Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		Non conformance / must state in issue if its an observation						
<a href="#">359497</a> <a href="#">25 Mar 2025</a>	Review Paper Customer Complaints File (75)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
System Generated

Review Paper Customer Complaints File

**25 Mar 2025 Derek Lamb**  
reviewed in audit last week all upto date nothing new

Audit CE Files  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">359360</a> <a href="#">24 Mar 2025</a>	Audit 12 CE Files Viamed (16)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
System Generated BSI Audits Calendar BSI Audit CE Files Audit 12 BEFORE starting the Audit you need to update the Processes attached to the Audit. Find the doc in the Document Index Admin Document - Copy and Paste List Processes Per Title replacing them in the current audit. Review Last years Audit see if its still suitable Carry out the Audit, reviewing and commenting on the ISO route map first. If there are any non Conformances from the Audit, create a follow up / related Issue including all of the following - With a time for Completion Immediate Action Plan Corrective Action Plan Corrective Action Confirmation of Resolution If its a major / critical non conformance complete form QC 18

**31 Mar 2025 Helen Lamb**  
Next Action Changed From Helen Lamb To Derek Lamb audit 12 Technical files Viamed 2024 attached no outstanding issues please review

**01 Apr 2025 Derek Lamb**  
Done

Audit Organisation and Process Verification Internal Process Verification  
Viamed

Issue / Primary	Subject/Notes	Minor Internal Error i.e. not ISO	Reviewed Non Conformity	Determined Cause of Non	Evaluated action to Ensure	Planning and documenting action needed	Verify Action does not adversely	Effectiveness of corrective
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ID / Call ID		Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	/ Complaint and determine if its a vigilance Issue requiring a corrective action plan	Conformity / Complaint	does not recur	and implementation QC 28b	affect Safety Performance or regulatory requirements	action reviewed
<a href="#">358650</a> <a href="#">17 Mar 2025</a>	Email Distribution (366)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
System Generated Check the global mailbox, there should be no mails more than a Day old. Give allowance for Monday mornings, there should be no emails older than Friday afternoon. Also note size of space used in Global inbox

**17 Mar 2025 Derek Lamb**  
mail box empty

Supplier Complaints (us to them)  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">358580</a> <a href="#">14 Mar 2025</a>	test4	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
Added by Derek Lamb sent to Derek Lamb  
test4

**14 Mar 2025 Derek Lamb**

Supplier Complaints (Them to Us)  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">358579</a> <a href="#">14 Mar 2025</a>	test3	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
Added by Derek Lamb sent to Derek Lamb  
test3

**14 Mar 2025 Derek Lamb**

Complaints from VST to a Supplier  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">358578</a> <a href="#">14 Mar 2025</a>	test2	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>



		in issue if its an observation						
<a href="#">358213</a> <a href="#">11 Mar</a> <a href="#">2025</a>	Order Error : 155158 Viamed Memo error	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb  
Auto Issue from Error Log 155158  
Order Entered by Kate Griffiths  
Order Checked by Sophie Lines  
Goods Out  
Error was Viamed  
Memo error  
Fault:  
Serial numbers too long for the invoice

Possible Fix  
Suggested only have 2 sets of matched pairs per line.

**Audit Goods Inwards and Product Identity**  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">357925</a> <a href="#">10 Mar</a> <a href="#">2025</a>	Audit 09 Goods Inward And Product Identity Viamed (170)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
System Generated Audit 09 BEFORE starting the Audit you need to update the Processes attached to the Audit. Find the doc in the Document Index Admin Document - Copy and Paste List Processes Per Title replacing them in the current audit. Review Last years Audit see if its still suitable Carry out the Audit, reviewing and commenting on the ISO route map first. If there are any non Conformances from the Audit, create a follow up / related Issue including all of the following - With a time for Completion Immediate Action Plan Corrective Action Plan Corrective Action Confirmation of Resolution If its a major / critical non conformance complete form QC 18

**09 May 2025 Helen Lamb**  
Next Action Changed From Helen Lamb To Derek Lamb completed audit attached. Nothing ongoing no problems or issues. nothing from last audit or non conformances. Please review

**09 May 2025 Derek Lamb**  
thankyou

**Non Conformance Issues**

Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">357750</a> <a href="#">06 Mar 2025</a>	Shipped Items Return to Supplier BOX1023	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
Does this Return BOX1023 warrant a NON conformance report via the CAPA process VM3COP10

**07 Mar 2025 Derek Lamb**  
for calibration no issues

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">357468</a> <a href="#">05 Mar 2025</a>	Non conformance review history Viamed (283)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
System Generated Check the history of the last Non conformance review, check actions are being carried out, and non conformances are not reoccurring. Review the Order Invoice Error Logs (link in issue) for increases in errors. Areas that are errors that are our mistake and not due to changes instigated by the customer. Note any problem areas in report. Review Customer Complaint and Non Conformance Review Screen check back over the previous month since last review to see if

anything internal / staff / process related needs to be highlighted. Go to Add an issue, search non con and review any issues since the last review for internal / staff / process related issues that needs to be highlighted. Note any problems in the issue and send to Derek to review

**17 Mar 2025 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb Checked back through all Non Conformance issues (not automatically generated) to 1st March 25. Issue 354351 (also 354400) high failure rate on 0110361. 355521 Ceratherm 600-2 and 600-3 Casing. holes drilled in incorrect place, replaced by supplier asked us to dispose of. 356739 MySign Ono cap required. 356891 cracked display. None of these require a capa. No problems. Nothing else of concern or to be investigated or reported on in Non Con issues. Review VIAMED Feedback - Product Feedback Negative (741) nothing new. Review VIAMED Feedback - Product Feedback Positive (1189) nothing new. Review VIAMED Feedback - Customer Feedback Negative (739) nothing new. Review VIAMED Feedback - Customer Feedback Positive (1188) nothing new. Review VIAMED Feedback - Customer Complaints (737) nothing new. Order Invoice Error Logs - a lot of address amends but some are customer and only a few are input error based. Some spelling errors. Will speak to the office about being more careful. No issue, nothing to worry about. Non Conformance, complaints and feedback headers nothing else and nothing of concerns. Test issues have been sent to make sure they show in the correct areas. Several issues in the non conformance header re error logging. All checked and where needed relevant staff have been spoken to. Non Conformance issues review screen - No issues not already reviewed above in log. No other feedback issues relating to non conformances that need to be monitored or reviewed. Derek please review the above

**17 Mar 2025 Derek Lamb**

thankyou

**Audit Analysis of Data**  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">357464</a> <a href="#">05 Mar 2025</a>	Review Company Data (114)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

**Derek Lamb**

System Generated Review Sales - Number Invoices / Credits per day reports Search for any potential problems in the Data provided

Employee Menu Check Audit Roles Titles and Processes in Employee Roles and Titles. Check the Page for Red Crosses and potentially missing objectives Search the Outstanding Jobs column for potential problems Do we have enough resources. Check the Issues Statistics in the meeting header, Check Issues are being completed in a timely manner.

**10 Mar 2025 Derek Lamb**

reviewed, went through issues roles with office last wednesday

**Non Conformance Issues**  
Viamed

Issue / Primary	Subject/Notes	Minor Internal Error	Reviewed Non	Determined Cause of	Evaluated action to	Planning and documenting	Verify Action does not	Effectiveness of corrective
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<a href="#">2025</a>	Missing info - office							
<p>Helen Lamb          Auto Issue from Error Log 155201          Order Entered by Emily Hanson          Order Checked by Aqib Majeed          Office          Error was Viamed          Missing info - office          Fault:          wrong quantity was for 3 we put 2 so had wrong price. Credited difference and sent third posey box at the 3 off price.</p> <p>Possible Fix</p>								

**Non Conformance Issues**  
 Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">357181</a> <a href="#">03 Mar 2025</a>	0110361 QA Data Requires Risk / Non Conformance Review Feb 2025	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
 Added by Robert Connor sent to Derek Lamb  
 INFORMATION ONLY ISSUE **DO NOT** ADD NOTES!  
 QA Failures High Numbers

**Audit Documentation Control**  
 Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested /	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation						
<a href="#">357077</a> <a href="#">03 Mar 2025</a>	Documentation out of date (372)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb  
System Generated  
Check for Out of Date documents

This is an audit, you do not need to perform the Task:

Simply ensure all out of date documents have an Issue attached to get them updated.  
If the Issue is more than 2 Months out of date read the issue - if appropriate generate a non conformance Issue

ISO - Document index admin

Scroll down and check if any documents have gone out of date,

Either update the document or create an Issue to the relevant person from the document admin / details screen.  
Remember if you update a document reset the expiry date

**03 Mar 2025 Helen Lamb**  
Done

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">356891</a> <a href="#">27 Feb 2025</a>	Shipped Items Return to Supplier BOX1021	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
Does this Return BOX1021 warrant a NON conformance report via the CAPA process VM3COP10

**28 Feb 2025 Derek Lamb**  
The unit has a cracked display physical damage, no cap at this time 2510091 tof

Non Conformance Issues Viamed								
Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">356876</a> <a href="#">27 Feb</a> <a href="#">2025</a>	Order Error : 155251 Viamed ebay error - goods out	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb  
Auto Issue from Error Log 155251  
Order Entered by Kate Griffiths  
Order Checked by Emily Hanson  
Non Selected  
Error was Viamed  
ebay error - goods out  
Fault:  
Order not fulfilled on eBay

Possible Fix  
Done

Non Conformance Issues Viamed								
Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed



		actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Issue requiring a corrective action plan					
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<a href="#">356467</a> <a href="#">24 Feb</a> <a href="#">2025</a>	Order Error : 153642 Viamed Memo error	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Helen Lamb  
Auto Issue from Error Log 153642  
Order Entered by Aqib Majeed  
Order Checked by Sophie Lines  
Director  
Error was Viamed  
Memo error  
Fault:  
TEST ERROR NOT REAL

Possible Fix

Audit Contract Review  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">355841</a> <a href="#">18 Feb</a> <a href="#">2025</a>	Distributor Agreements (379)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
System Generated  
Note this is an Audit - simply need to ensure its being carried out

Sales -> Distributor Agreements -> Check Sales Against Agreements,

List should be up to date / empty.

**20 Feb 2025 Derek Lamb**  
task appears reasonably upto date

**Audit Organisation and Process Verification Internal Process Verification**  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">355681</a> <a href="#">17 Feb 2025</a>	Email Distribution (366)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb

System Generated Check the global mailbox, there should be no mails more than a Day old. Give allowance for Monday mornings, there should be no emails older than Friday afternoon. Also note size of space used in Global inbox

**20 Feb 2025 Derek Lamb**  
reviewed

**Non Conformance Issues**  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		Non conformance / must state in issue if its an observation						
<a href="#">355495</a> <a href="#">13 Feb</a> <a href="#">2025</a>	Order Error : 153642 Missing info - office	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb  
Auto Issue from Error Log 153642  
Order Entered by Aqib Majeed  
Order Checked by Sophie Lines  
Office  
Error was Missing info - office  
Fault:  
Incorrect Inco term used on a EXW order

Possible Fix  
passed to helen to correct

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">355144</a> <a href="#">10 Feb</a> <a href="#">2025</a>	Order Error : 154818 wrong account - office	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Helen Lamb  
Auto Issue from Error Log 154818  
Order Entered by Aqib Majeed  
Order Checked by Kate Griffiths  
Office  
Error was wrong account - office  
Fault:

Possible Fix

Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">355055</a> <a href="#">10 Feb 2025</a>	Audit 27 Software Validation Viamed (821)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb

System Generated Task To be Completed Audit 27 Software Validation To confirm the Prime functions of the Software used is verified. The Audit itself, Intrastats, physical process being carried out. Complete Audit 27 BEFORE starting the Audit you need to update the Processes attached to the Audit. Find the doc in the Document Index Admin Document - Copy and Paste List Processes Per Title replacing them in the current audit. Review Last years Audit see if its still suitable Carry out the Audit, reviewing and commenting on the ISO route map first. If there are any non Conformances from the Audit, create a follow up / related Issue including all of the following - With a time for Completion Immediate Action Plan Corrective Action Plan Corrective Action Confirmation of Resolution If its a major / critical non conformance complete form QC 18

**11 Feb 2025 Helen Lamb**

completed audit attached, no issues or problems. Nothing outstanding.

Non Conformance Issues

Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">355001</a> <a href="#">07 Feb</a>	Shipped Items Return to Supplier BOX1016	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

2025

Derek Lamb  
Does this Return BOX1016 warrant a NON conformance report via the CAPA process VM3COP10

**10 Feb 2025 Derek Lamb**  
0110432History of 25464842546484 KF69099174Non Linear normal sensor type, no cap required

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">354707</a> <a href="#">05 Feb 2025</a>	Shipped Items Return to Supplier BOX1017	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
Does this Return BOX1017 warrant a NON conformance report via the CAPA process VM3COP10

**06 Feb 2025 Derek Lamb**  
calibration no cap

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		an observation						
<a href="#">354618</a> <a href="#">05 Feb</a> <a href="#">2025</a>	Non conformance review history Viamed (283)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb  
System Generated Check the history of the last Non conformance review, check actions are being carried out, and non conformances are not reoccurring. Review the Order Invoice Error Logs (link in issue) for increases in errors. Areas that are errors that are our mistake and not due to changes instigated by the customer. Note any problem areas in report. Review Customer Complaint and Non Conformance Review Screen check back over the previous month since last review to see if anything internal / staff / process related needs to be highlighted. Go to Add an issue, search non con and review any issues since the last review for internal / staff / process related issues that needs to be highlighted. Note any problems in the issue and send to Derek to review

**10 Feb 2025 Helen Lamb**  
Checked back through all Non Conformance issues (not automatically generated) to 1st Jan 25. Issue 352880 wrong sensors in boxes still ongoing from last review. 354351 high failure rate on 0110361 R22AHJR sensors no cap. Nothing new to add, just resolving. Nothing else of concern or to be investigated or reported on in Non Con issues. Review VIAMED Feedback - Product Feedback Negative (741) nothing new. Review VIAMED Feedback - Product Feedback Positive (1189) nothing new. Review VIAMED Feedback - Customer Feedback Negative (739) nothing new. Review VIAMED Feedback - Customer Feedback Positive (1188) nothing new. Review VIAMED Feedback - Customer Complaints (737) nothing new. Order Invoice Error Logs - a lot of address amends but some are customer and only a few are input error based. No issue, nothing to worry about. Non Conformance, complaints and feedback headers 351507 a request for PMS, 353625 Ceratherm cases incorrectly drilled - ongoing, 351010 + 354400 0110361 QA Data Requires Risk / Non Conformance Review Dec 2024, 351194 + 351875Incorrect oxygen sensors in boxes, 353250, 353881, 354144, 354707, 355001 auto issues. Everything underway nothing else and nothing of concerns. Non Conformance issues review screen - No issues not already reviewed above in log. No other feedback issues relating to non conformances that need to be monitored or reviewed. Derek please review the above

**Audit Analysis of Data**  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed

<a href="#">354614</a> <a href="#">05 Feb</a> <a href="#">2025</a>	Review Company Data (114)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
System Generated Review Sales - Number Invoices / Credits per day reports Search for any potential problems in the Data provided  
  
Employee Menu Check Audit Roles Titles and Processes in Employee Roles and Titles. Check the Page for Red Crosses and potentially missing objectives Search the Outstanding Jobs column for potential problems Do we have enough resources. Check the Issues Statistics in the meeting header, Check Issues are being completed in a timely manner.

**06 Feb 2025 Derek Lamb**  
reviewed



Derek Lamb  
 Added by Derek Lamb sent to Derek Lamb  
 Oxygen sensor R-22AHJR Jikco OEM sensor known yield, no cap required

**03 Feb 2025 Derek Lamb**

**21 Feb 2025 Derek Lamb**

Non Conformance Issues  
 Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">354400</a> <a href="#">03 Feb 2025</a>	0110361 QA Data Requires Risk / Non Conformance Review Jan 2025	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
 Added by Robert Connor sent to Derek Lamb  
 INFORMATION ONLY ISSUE **DO NOT** ADD NOTES!  
 QA Failures High Numbers

**03 Feb 2025 Derek Lamb**

Created Related Issue #354461  
 Added by Derek Lamb sent to Derek Lamb  
 Oxygen sensor R-22AHJR Jikco OEM sensor known yield, no cap required

Non Conformance Issues  
 Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		no requirement to Escalate Non conformance / must state in issue if its an observation						
<a href="#">354239</a> <a href="#">31 Jan 2025</a>	UnShipped Items Return to Supplier BOX956	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
Does this Return BOX956 warrant a NON conformance report via the CAPA process VM3COP10

**31 Jan 2025 Derek Lamb**  
automotive sensors normal sensor fail type, no cap required

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">354144</a> <a href="#">30 Jan 2025</a>	Order Error : 153234 wrong account - office	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb  
Auto Issue from Error Log 153234  
Order Entered by Emily Hanson  
Order Checked by Aqib Majeed  
Office  
Error was wrong account - office  
Fault:  
on wrong account should have been Scottish Ambulance Service. also no email address for invoices either

Possible Fix  
better checking and looking at paperwork

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non	Reviewed Non Conformity /	Determined Cause of Non	Evaluated action to Ensure	Planning and documenting action needed and	Verify Action does not adversely affect Safety	Effectiveness of corrective action reviewed
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		Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Complaint and determine if its a vigilance Issue requiring a corrective action plan	Conformity / Complaint	does not recur	implementation QC 28b	Performance or regulatory requirements	
<a href="#">353881</a> <a href="#">28 Jan 2025</a>	Order Error : 154654 ebay error - goods out	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb  
Auto Issue from Error Log 154654  
Order Entered by Kate Griffiths  
Order Checked by Emily Hanson  
Goods Out  
Error was ebay error - goods out  
Fault:  
Order not fulfilled on Ebay

Possible Fix  
Done

Complaints from Viamed to a Supplier  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">353625</a> <a href="#">24 Jan 2025</a>	Ceratherm cases incorrectly drilled	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Catherine Spence  
Added by Steve Hardaker sent to Catherine Spence

Burnley General has reported that they received a Ceratherm case 0330000 that has holes that are incorrectly drilled, the front one is higher than the back one. Please can you check al existing stock ASAP as we need to send a replacement to them, see SRS69010.

**24 Jan 2025 Catherine Spence**

done, replacement sent

**24 Jan 2025 Steve Hardaker**

They need a returns label sending to collect the faulty one. Please can you generate a returns label for the faulty Ceratherm case on SRS69010, the address is as follows: Clinical Engineering Department Burnley General Hospital Casterton Avenue, Burnley, Lancashire. BB10 2PY Billy Rafiq - tel: 01254 733140

**27 Jan 2025 Catherine Spence**

UPS Label generated and passed to office to send to customer

**31 Jan 2025 Steve Hardaker**

Next Action Changed From Catherine Spence To General Sent pictures to Daniel at Nufer and asked if they want it sending back.

**03 Feb 2025 Steve Hardaker**

Next Action Changed From General To Catherine Spence Ewimed Nufer have agreed to send a replacement FOC and said to not bother sending the faulty one back. Not sure if this will need some paperwork to book it?

**13 Feb 2025 Catherine Spence**

will generate the return so that we can accept the replacement in to stock. No stock will be returned to nufer

**13 Feb 2025 Steve Hardaker**

Urgent Flag Changed To Off When you have the resolved returns reference from Nufer, can you add it here for closure, thanks.

**13 Feb 2025 Catherine Spence**

RMA was created for internal purpose only Box 1019, will complete off with replacement already received from supplier

**13 Feb 2025 Steve Hardaker**

Viamed Audits Calander  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">353284</a> <a href="#">22 Jan 2025</a>	happy to continue schedule	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

General  
22 Jan 2025 Derek Lamb

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">353250</a> <a href="#">22 Jan 2025</a>	Order Error : 154568 New Error	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb  
Auto Issue from Error Log 154568  
Order Entered by Sophie Lines  
Order Checked by Kate Griffiths  
Goods Out  
Error was New Error  
Fault:  
Order not fulfilled on Shopify

Possible Fix  
Done

Audit Organisation and Process Verification Internal Process Verification  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		no requirement to Escalate Non conformance / must state in issue if its an observation						
<a href="#">352212</a> <a href="#">10 Jan 2025</a>	Shipped Items Return to Supplier BOX1012	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
Does this Return BOX1012 warrant a NON conformance report via the CAPA process VM3COP10

**10 Jan 2025 Derek Lamb**  
8010006 (0.32 %) (vst) normal o2 senor fail type no cap required

Future Reviews - Internal Audits  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">351914</a> <a href="#">08 Jan 2025</a>	Objective Ensure the Audits are performed within a timely manner Review the Tasks and Audits for the Audits Should be no more than 1 outstanding issue for each section	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

General  
08 Jan 2025 Derek Lamb  
few backuoed due to christmas, but realtively oke

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective	Reviewed Non Conformity / Complaint and determine if its a vigilance	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Issue requiring a corrective action plan						
<a href="#">351875</a> <a href="#">08 Jan 2025</a> 351194	QC 21 Non Conformance Report - Incorrect oxygen sensors in boxes Non Conformance	<input type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>

Derek Lamb  
 Added by Helen Lamb sent to Derek Lamb  
 From issue 351194. There have been some oxygen sensors that have been boxed, but have been placed in the wrong box with the wrong barcode attached. Original issue from Cathy 3rd January 25. To be investigated.

**24 Feb 2025 Helen Lamb** this has been resolved and no reoccurrence has happened. Staff have been re trained and a new policy where no different types of stock as placed together for boxing has been implemented. QC21 form needs to be updated

**26 Feb 2025 Helen Lamb** this has been resolved and no reoccurrence has happened. Staff have been re trained and a new policy where no different types of stock as placed together for boxing has been implemented. QC21 form needs to be updated

Non Conformance Issues  
 Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">351232</a> <a href="#">06 Jan 2025</a>	Non conformance review history Viamed (283)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
 System Generated Check the history of the last Non conformance review, check actions are being carried out, and non conformances are not reoccurring. Review the Order Invoice Error Logs (link in issue) for increases in errors. Areas that are errors that are our mistake and not due to changes instigated by the customer. Note any problem areas in report. Review Customer Complaint and Non Conformance Review Screen check back over the previous month since last review to see if anything internal / staff / process related needs to be highlighted. Go to Add an issue, search non con and review any issues since the last review for internal / staff / process related issues that needs to be highlighted. Note any problems in the issue and send to Derek to review

**08 Jan 2025 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb Checked back through all Non Conformance issues (not automatically generated) to 1st Dec 24. 350995 0110361 R22AHJR high failure rate. Nothing on going. Nothing else of concern or to be investigated or reported on in Non Con issues. Review VIAMED Feedback - Product Feedback Negative (741) nothing new. Review VIAMED Feedback - Product Feedback Positive (1189) nothing new. Review VIAMED Feedback - Customer Feedback Negative (739) nothing new. Review VIAMED Feedback - Customer Feedback Positive (1188) nothing new. Review VIAMED Feedback - Customer Complaints (737) nothing new. Order Invoice Error Logs - no issue, nothing to worry about. Nothing else new, no concerns. Non Conformance, complaints and feedback headers reviewed, Issue 351194 Wrong sensors placed in wrong boxes and have been sold QC21 form to be generate. Non Conformance issues review screen - No issues not already reviewed above in log. No other feedback issues relating to non conformances that need to be monitored or reviewed. Derek please review the above

**08 Jan 2025 Derek Lamb**

thankyou

**Audit Analysis of Data**  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">351223</a> <a href="#">06 Jan 2025</a>	Review Company Data (114)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb

System Generated Review Sales - Number Invoices / Credits per day reports Search for any potential problems in the Data provided

Employee Menu Check Audit Roles Titles and Processes in Employee Roles and Titles. Check the Page for Red Crosses and potentially missing objectives Search the Outstanding Jobs column for potential problems Do we have enough resources. Check the Issues Statistics in the meeting header, Check Issues are being completed in a timely manner.

**06 Jan 2025 Derek Lamb**

just been christmas so lots of staff off, not a suitable time to review outstanding issues, they need time to catch up,

**Non Conformance Issues**  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective	Reviewed Non Conformity / Complaint and determine if its a vigilance	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Issue requiring a corrective action plan					
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<a href="#">351203</a> <a href="#">03 Jan</a> <a href="#">2025</a>	UnShipped Items Return to Supplier BOX986	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
Does this Return BOX986 warrant a NON conformance report via the CAPA process VM3COP10

**06 Jan 2025 Derek Lamb**  
nitrogen sensor leaked, normal fail type, single unit, no cap required

Non Conformance Issues  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">351194</a> <a href="#">03 Jan</a> <a href="#">2025</a>	Incorrect oxygen sensors in boxes	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Helen Lamb  
Added by Catherine Spence sent to Helen Lamb  
There have been some oxygen sensors that have been boxed, but have been placed in the wrong box with the wrong barcode attached. R-49V (0110049) S/N: V104581, one of these was boxed, but inside the box was a T7V oxygen sensor S/N V101681 RVM153007-1 25/10/24 we have replaced the incorrect sensor on RVM154184-1 and made SRS68990 for the return of the incorrect sensor and also sent out a prepaid UPS label for the return. 12 x 0110022 sold to Broomfield Hosp on RVM153914-1 on 13/12/24 we have been informed that four of them were in fact R-17meds 0110017 We have replaced the full order on RVM154163-1 and have generated SRS68989 for the return of the sensors, and supplied a UPS prepaid returns label. The effected s/n of the R17Meds are which in turn will now have R-22med sensors in side are. S/N:165021 Leeds Hosp RVM15394-1 dated 17/12/23, replacement sensor sent out today on RVM154192-1 FOC S/N:165022 Morrison Hosp RVM154065-1 dated 20/12/24, replacement sent out today on RVM154189-1 FOC S/N:165023 ESZ AG Calibration RVM154064-1 dated 23/12/24, replacement sent out on RVM154188-1 FOC S/N:165024 Royal Alexander Hosp RVM154073-1 dated 23/12/24, replacement sent out today RVM154185-1 FOC All effected orders have been informed of the issue. It would appear that the boxer has only checked the last two digits of the sensor and not carried out a full check to match the type of sensor and whole serial number from the unit to the barcode label. All staff who have access to boxing of sensors will

be sent an issue to what has happened, and all staff will be re trained. All current pre boxed sensors that are on the ready for sale shelf are being re checked before any other orders can go out. Going forward, we will limit the amount of people able to box sensors to : Linda, myself and robert. And for the moment, before we box any sensors we will scan then to our own location , so that we can confirm who was boxing sensors.( Linda`s sensors will be pre scanned to her location for her) once all boxing has been done then they will be scanned to sellable shelf.

**08 Jan 2025 Helen Lamb**

Created Related Issue #351875

Added by Helen Lamb sent to Derek Lamb

From issue 351194. There have been some oxygen sensors that have been boxed, but have been placed in the wrong box with the wrong barcode attached. Original issue from Cathy 3rd January 25. To be investigated.

**16 Jan 2025 Helen Lamb**

RVM154184 is seaward and sensor has been returned. Credit done TVM153007-1

**24 Feb 2025 Helen Lamb**

this has been resolved and no reoccurrence has happened. Staff have been re trained and a new policy where no different types of stock as placed together for boxing has been implemented.

**14 May 2025 Catherine Spence**

**Non Conformance Issues**

Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">351010</a> <a href="#">02 Jan 2025</a>	0110361 QA Data Requires Risk / Non Conformance Review Dec 2024	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb

Added by Robert Connor sent to Derek Lamb

INFORMATION ONLY ISSUE **DO NOT** ADD NOTES!

QA Failures High Numbers

**Audit Design Control**

Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective	Reviewed Non Conformity / Complaint and determine if its a vigilance	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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		actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Issue requiring a corrective action plan					
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<a href="#">350895</a> <a href="#">01 Jan</a> <a href="#">2025</a>	Audit 03 Design Control VST (193)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
System Generated Audit 03 NOTE DESIGN REMOVED FROM VST, AUDIT NOT REQUIRED. LEFT IN FOR FUTURE USE Review Last years Audit see if its still suitable Before Proceeding you need to update the Processes attached to the Audit Search the Document in the Index, View the Admin Page Copy and Paste the Attached Processes, replacing them in the current audit Any non Conformances from the Audit: Create a follow up / related Issue, With a time for Completion Immediate Action Plan Corrective Action Plan Corrective Action Confirmation of Resolution if its a major / critical non conformance complete form QC 18

**22 Jan 2025 Helen Lamb**  
signed off in error in meeting closing 2024. Audit carried out 22nd Jan 25

**22 Jan 2025 Helen Lamb**  
Next Action Changed From Helen Lamb To Derek Lamb Audit completed no issue ongoing no problems

**23 Jan 2025 Derek Lamb**  
No requiremnet for this audit anymore, design sections not included in the ISO

Audit Picking and Packing  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
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<a href="#">350890</a> <a href="#">01 Jan</a> <a href="#">2025</a>	Audit 01 Picking Packing Viamed (24)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
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Derek Lamb  
System Generated BEFORE starting Audit you need to update the Processes attached to the Audit. Find the doc in the Document Index Admin Document - Copy and Paste List Processes Per Title replacing them in the current audit. Review Last

years Audit see if its still suitable Carry out the Audit, reviewing and commenting on the ISO route map first. If there are any non Conformances from the Audit, create a follow up / related Issue including all of the following - With a time for Completion Immediate Action Plan Corrective Action Plan Corrective Action Confirmation of Resolution If its a major / critical non conformance complete form QC 18

**22 Jan 2025 Helen Lamb**

signed off in error in meeting closing 2024. Audit carried out 22nd Jan 25

**27 Jan 2025 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb Issue completed and no problems or outstanding issues.

**27 Jan 2025 Derek Lamb**

reviewed, thankyou

**Audit Design Control**  
Viamed

Issue / Primary ID / Call ID	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / must state in issue if its an observation	Reviewed Non Conformity / Complaint and determine if its a vigilance Issue requiring a corrective action plan	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure does not recur	Planning and documenting action needed and implementation QC 28b	Verify Action does not adversely affect Safety Performance or regulatory requirements	Effectiveness of corrective action reviewed
<a href="#">350889</a> <a href="#">01 Jan 2025</a>	Audit 03 Design Control Viamed (22)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

**Derek Lamb**

Audit 03 Design Control Review Last years Audit see if its still suitable Before Proceeding you need to update the Processes attached to the Audit Search the Document in the Index, View the Admin Page Copy and Paste the Attached Processes, replacing them in the current audit Any non Conformances from the Audit: Create a follow up / related Issue, With a time for Completion Immediate Action Plan Corrective Action Plan Corrective Action Confirmation of Resolution if its a major / critical non conformance complete form QC 18

**22 Jan 2025 Helen Lamb**

signed off in error in meeting closing 2024. Audit carried out 22nd Jan 25

**22 Jan 2025 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb completed audit attached no issues nothing outstanding.

**23 Jan 2025 Derek Lamb**

Design removed formt he ISO standards, audit not really required any more