

EH Full



## Customer Complaint and Non Conformance Review Screen

[Show All Generic Issues](#)  
[Show All Viamed Issues](#)  
[Show All Vandagraph Issues](#)  
[Show All VST Issues](#)  
[Show All Human Med Issues](#)  
[Show All Viamed Properties Issues](#)  
[Show All The Pointless Logo Company Issues](#)  
[Show All Non Minor Issues](#)

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

Non Conformance Issues								
Viamed Vandagraph VST Human Med 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 / or dealt with in s	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="#">351233</a> <a href="#">06 Jan 2025</a>	Non conformance review history VST (285)	<input 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								
Non Conformance Issues								
Viamed Vandagraph VST Human Med 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 / or dealt with in s	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 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								
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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 / or dealt with in s	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="#">351194</a> <a href="#">03 Jan 2025</a>	Incorrect oxygen sensors in boxes	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

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.

## Audit Picking and Packing

Viamed Vandagraph VST Human Med 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 / or dealt with in s	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="#">350896</a> <a href="#">01 Jan 2025</a>	Audit 01 Picking Packing VST (194)	<input 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 01 due 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

## Audit Design Control

Viamed Vandagraph VST Human Med 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 / or dealt with in s	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="#">350895</a> <a href="#">01 Jan 2025</a>	Audit 03 Design Control VST (193)	<input 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 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

## Audit Picking and Packing

Viamed Vandagraph VST Human Med 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 / or dealt with in s	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="#">350890</a> <a href="#">01 Jan 2025</a>	Audit 01 Picking Packing Viamed (24)	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen 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

## Audit Design Control

Viamed Vandagraph VST Human Med 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 / or dealt with in s	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 type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen 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

## Non Conformance Issues

Viamed Vandagraph VST Human Med 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 / or dealt with in s	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="#">350765</a> <a href="#">30 Dec 2024</a>	Order Error : 154112 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 154112

Order Entered by Aqib Majeed

Order Checked by Emily Hanson

Office

Error was Checking error

Fault:

Has been entered on the wrong priority, should be a pri 4 as its royal mail

Possible Fix

passed back to the office

## Non Conformance Issues

VST

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<a href="#">349901</a> <a href="#">17 Dec 2024</a>	Shipped Items Return to Supplier BOX1010	<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 BOX1010 warrant a NON conformance report via the CAPA process VM3COP10

**17 Dec 2024 Derek Lamb**

vst sensors normal fail type no capa required

## Non Conformance Issues

VST

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<a href="#">348823</a> <a href="#">05 Dec 2024</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

**13 Dec 2024 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb Checked back through all Non Conformance issues (not automatically generated) to 1st Nov 24. 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 issue, nothing to worry about. Non Conformance, complaints and feedback headers reviewed positive feedback received, 344745 positive Feedback from Air Research 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 Dec 2024 Derek Lamb**

thankyou

**Audit of Audits**

Viamed Vandagraph VST Human Med 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 / or dealt with in s	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="#">348378</a> <a href="#">02 Dec 2024</a>	Audit 21 Audit Of Audit VST (192)	<input 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 21 due Review the Audit Calendar Screen ISO -> Audit Calendar Complete Audit 21 Confirm if Audit calendar needs changing. 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

**Audit Internal Audits**

Viamed Vandagraph VST Human Med 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 / or dealt with in s	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="#">348377</a> <a href="#">02 Dec 2024</a>	Audit 17 Internal Audits VST (191)	<input 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 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. 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

**Audit Internal Audits**

Viamed Vandagraph VST Human Med Viamed Properties The Pointless Logo Company

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<a href="#">348369</a> <a href="#">02 Dec 2024</a>	Audit 17 Internal Audits Viamed (11)	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb

System Generated BSI Audits Calendar BSI Audit Internal Audits 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. 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

Audit Customer Complaints VST								
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 / or dealt with in s	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="#">346837</a> <a href="#">14 Nov 2024</a>	Audit 14 Complaints And Corrective Actions VST (189)	<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 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. 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

**18 Nov 2024 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb completed audit attached please review. Nothing outstanding

**18 Nov 2024 Derek Lamb**

thankyou

## BSI Minor Non conformances

[Viamed](#) [Vandagraph](#) [VST](#) [Human Med](#) [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 / or dealt with in s	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="#">346434</a> <a href="#">08 Nov 2024</a> 346177	2560663-202410-N1 Calibration Stickers	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Philip Crossley

Added by Derek Lamb sent to Philip Crossley

CE Number Barcode Serial Number Date Due CE058 102042 50613890 Dec/2025 CE059 102043 50613891 Dec/2025 CE076 102060 6939K9 Dec/2024 CE078 102062 04314 Jul/2025 CE082 102066 Unknown Oct/2026 CE138 102122 Z28-0458 Nov/2024 CE109 102094 D19477292 Sep/2018 CE111 102095 D19477299 Oct/2026 CE112 102096 D19477379 Apr/2025 CE149 102133 420720660 Jul/2025 CE169 102153 N/K Dec/2025 CE178 674700 530569434 Jul/2025 CE182 697263 2433026 Jul/2025 CE185 843663 425571 Oct/2025 CE189 995710 Unknown Oct/2026 CE195 996365 Unknown Oct/2026 CE206 1288063 3027310 Aug/2025 CE207 1303613 BM82699024 Sep/2025 CE214 2318147 Q232100069 Sep/2024 CE215 2325084 2059802/11 Oct/2025 CE216 2357823 578952 Dec/2024 CE218 2578411 Q231600080 Sep/2025 CE220 2591710 BX1651896 Nov/2028 CE221 2610182 2077396 Oct/2025 List of equipment to find to put stickers on

**11 Nov 2024 Philip Crossley**

Ok, I will print the list out and work through it

**11 Nov 2024 Philip Crossley**

Done - almost. 2 things are at Lindas - CE082 and CE111. I will pass stickers to Mike. 2 things are away for external calibration. CE076 Microcal and CE138 Seaward pat tester. They will be stickered when they return. 2 things are not in service - CE182 and CE214. I still need to find CE109, a Caltex DVM that is indication only.

**11 Nov 2024 Philip Crossley**

There are things missing from the list already done - CE091 CE094 maybe CE0179 CE198 CE210 CE219

**12 Nov 2024 Philip Crossley**

Added Excel of Calibration Sticker status with notes etc

**13 Nov 2024 Philip Crossley**

CE109 is lost, not seen since 2018 so I have removed it from service. I think the list is completed so I will close the issue.

## Non Conformance Issues

VST								
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<a href="#">346424</a> <a href="#">08 Nov 2024</a>	UnShipped Items Return to Supplier BOX1005	<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 BOX1005 warrant a NON conformance report via the CAPA process VM3COP10

**11 Nov 2024 Derek Lamb**  
automotive sensors, not being returned but replace by teledyne. no capa required

#### BSI Minor Non conformances

<div> <div>Viamed</div> <div>Vandagraph</div> <div>VST</div> <div>Human Med</div> <div>Viamed Properties</div> <div>The Pointless Logo Company</div> </div>								
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<a href="#">346177</a> <a href="#">06 Nov 2024</a>	2560663-202410-N1 Calibration Stickers	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Helen Lamb  
Added by Derek Lamb sent to Helen Lamb  
7.6 ISO 13485 Monitoring and measuring equipment process was not fully effective for servicing activities due to calibrated equipment was not appropriately labelled with current Calibration status nor identifiable.

**06 Nov 2024 Helen Lamb**  
first draft attached also training doc re calibration test equipment first draft

**08 Nov 2024 Derek Lamb**  
Created Related Issue #346434  
Added by Derek Lamb sent to Philip Crossley  
CE Number Barcode Serial Number Date Due CE058 102042 50613890 Dec/2025 CE059 102043 50613891 Dec/2025 CE076 102060 6939K9 Dec/2024 CE078 102062 04314 Jul/2025 CE082 102066 Unknown Oct/2026 CE138 102122 Z28-0458 Nov/2024 CE109 102094 D19477292 Sep/2018 CE111 102095 D19477299 Oct/2026 CE112 102096 D19477379 Apr/2025 CE149 102133 420720660 Jul/2025 CE169 102153 N/K Dec/2025 CE178 674700 530569434 Jul/2025 CE182 697263 2433026 Jul/2025 CE185 843663 425571 Oct/2025 CE189 995710 Unknown Oct/2026 CE195 996365 Unknown Oct/2026 CE206 1288063 3027310 Aug/2025 CE207 1303613 BM82699024 Sep/2025 CE214 2318147 Q232100069 Sep/2024 CE215 2325084 2059802/11 Oct/2025 CE216 2357823 578952 Dec/2024 CE218 2578411 Q231600080 Sep/2025 CE220 2591710 BX1651896 Nov/2028 CE221 2610182 2077396 Oct/2025 List of equipment to find to put stickers on

**12 Nov 2024 Derek Lamb** See Issue 346434, Phil has added stickers to existing Equipment, all bat the items currently being sent off for routine calibration, and 1 Sticker to be sent to Lindas test equipment.

**12 Nov 2024 Helen Lamb**  
update doc to include issue number and task ids updated

**12 Nov 2024 Helen Lamb**

#### Non Conformance Issues VST

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 / or dealt with in s	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="#">345968</a> <a href="#">05 Nov 2024</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

**07 Nov 2024 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb Checked back through all Non Conformance issues (not automatically generated) to 1st Oct 24. Issue 345034 Supplier return. Nothing else 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 issue, nothing to worry about. Non Conformance, complaints and feedback headers reviewed positive feedback received, 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

**12 Nov 2024 Derek Lamb**  
thank you

Audit Organisation and Process Verification  
VST

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 / or dealt with in s	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="#">345560</a> <a href="#">01 Nov 2024</a>	Audit 20 Process Verification To Managment VST (181)	<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 20 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 Nov 2024 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb Audit completed issue 347600 still ongoing re adding HSE sections to processes pages. Please review

**22 Nov 2024 Derek Lamb**

thankyou

Non Conformance Issues  
VST

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 / or dealt with in s	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="#">345034</a> <a href="#">25 Oct 2024</a>	Shipped Items Return to Supplier BOX1004	<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 BOX1004 warrant a NON conformance report via the CAPA process VM3COP10

**25 Oct 2024 Derek Lamb**

Relates to original purchase order PST3675 booked in 01-08-2024, sensors dated 2024-08. Envitec invoice: 5267472995. Original problem was due to linearity error at high pp02 &gt; 2.0 bar.

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 / or dealt with in s	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="#">345017</a> <a href="#">25 Oct 2024</a> 344455	Software Validation Document Control (802) Non Conformance	<input type="checkbox"/>	<input checked="" 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

document did not appear, page has been reviewed as to the reason , seems the filter for pulling out documents, that are not linked to a contact, failed, so we not showing the documents flagged as out of date, but not directly linked to a contact,. the bug has now been fixed, all documents are appearing,

**25 Oct 2024 Derek Lamb**

See QC 21 Form for document control.

Audit Management Review  
VST

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

		conformance / or dealt with in s						
<a href="#">343909</a> <a href="#">15 Oct 2024</a>	Audit 18 Management Review VST (188)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
<p>Helen Lamb</p> <p>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. 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</p>								
Non Conformance Issues VST								
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 / or dealt with in s	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="#">343426</a> <a href="#">09 Oct 2024</a>	Shipped Items Return to Supplier BOX998	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
<p>Derek Lamb</p> <p>Does this Return BOX998 warrant a NON conformance report via the CAPA process VM3COP10</p>								
<p><b>10 Oct 2024 Derek Lamb</b></p> <p>vst sensors normal fail type 8010006 Total Sales / Units In System : 45156 Warranty Fails : 118 0% Non Warranty Fails 37 0%</p>								
Non Conformance Issues VST								
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 / or dealt with in s	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="#">343025</a> <a href="#">07 Oct 2024</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"/>
<p>Derek Lamb</p> <p>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</p>								
<p><b>15 Oct 2024 Helen Lamb</b></p> <p>Next Action Changed From Helen Lamb To Derek Lamb Checked back through all Non Conformance issues (not automatically generated) to 1st Aug 24. 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 issue, nothing to worry about. Non Conformance, complaints and feedback headers reviewed 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</p>								
<p><b>16 Oct 2024 Derek Lamb</b></p> <p>thankyou</p>								
Non Conformance Issues VST								
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 / or dealt with in s	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="#">341623</a> <a href="#">21 Sep 2024</a>	Shipped Items Return to Supplier BOX993	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
<p>Derek Lamb</p> <p>Does this Return BOX993 warrant a NON conformance report via the CAPA process VM3COP10</p>								



**26 Sep 2024 Derek Lamb**  
vst jjcr normal oxygen sensor type fails

#### Non Conformance Issues VST

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 / or dealt with in s	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="#">340144</a> <a href="#">05 Sep 2024</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

#### 16 Sep 2024 Helen Lamb

Next Action Changed From Helen Lamb To Derek Lamb Checked back through all Non Conformance issues (not automatically generated) to 1st Aug 24. Nothing of concern or to be investigated or reported on. 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 issue, nothing to worry about. Non Conformance, complaints and feedback headers reviewed 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

#### 18 Sep 2024 Derek Lamb

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#### Audit Purchasing VST

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 / or dealt with in s	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="#">339679</a> <a href="#">02 Sep 2024</a>	Audit 05 Purchasing Suppliers VST (190)	<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

#### 17 Oct 2024 Helen Lamb

Next Action Changed From Helen Lamb To Derek Lamb completed audit attached. Nothing outstanding please review

#### 18 Oct 2024 Derek Lamb

thankyou

#### Audit Contract Review VST

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 / or dealt with in s	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="#">339678</a> <a href="#">02 Sep 2024</a>	Audit 02 Contract Review VST (187)	<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 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

**26 Sep 2024 Helen Lamb**  
 Created Related Issue #342152  
 Added by Helen Lamb sent to Derek Lamb  
 please find the updated audit attached, removed , no longer used task IDs re credit cards. We no longer write these out so no need to shred them

#### Non Conformance Issues VST

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 / or dealt with in s	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="#">339028</a> <a href="#">23 Aug 2024</a>	Shipped Items Return to Supplier BOX997	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Steve Nixon  
 Does this Return BOX997 warrant a NON conformance report via the CAPA process VM3COP10

#### 04 Sep 2024 Derek Lamb

Next Action Changed From Derek Lamb To Steve Nixon Dive soft, known issue: solved by Steve Nixon,

#### 30 Oct 2024 Steve Nixon

Priority Changed From 5 To 1

#### Non Conformance Issues VST

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 / or dealt with in s	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="#">338155</a> <a href="#">14 Aug 2024</a>	Shipped Items Return to Supplier BOX991	<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 BOX991 warrant a NON conformance report via the CAPA process VM3COP10

#### 14 Aug 2024 Derek Lamb

vst normal sensor fail types, no cap required

#### Non Conformance Issues VST

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 / or dealt with in s	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="#">337082</a> <a href="#">05 Aug 2024</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

#### 13 Aug 2024 Helen Lamb

Next Action Changed From Helen Lamb To Derek Lamb Checked back through all Non Conformance issues (not automatically generated) to 1st July 24. Nothing of concern or to be investigated or reported on. 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 issue, nothing to worry about. Non Conformance, complaints and feedback headers reviewed 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

#### 14 Aug 2024 Derek Lamb

thank you

### Audit Health and Safety VST

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 / or dealt with in s	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="#">336820</a> <a href="#">01 Aug 2024</a>	Audit 19 Health And Safety VST (186)	<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 Review Last years Audit see if its still suitable 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 Aug 2024 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb completed audit attached, nothing outstanding please review

**10 Sep 2024 Derek Lamb**

thankyou

### Audit Analysis of Data VST

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 / or dealt with in s	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="#">335184</a> <a href="#">15 Jul 2024</a>	Audit 23 Analysis Of Data VST (185)	<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 23 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

**16 Aug 2024 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb completed audit attached please review nothing ongoing

**10 Sep 2024 Derek Lamb**

thankyou

### Non Conformance Issues VST

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 / or dealt with in s	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="#">335086</a> <a href="#">12 Jul 2024</a>	Shipped Items Return to Supplier BOX979	<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 BOX979 warrant a NON conformance report via the CAPA process VM3COP10

**12 Jul 2024 Derek Lamb**

vst 8010020 normal sensor fail types no capa required

### Non Conformance Issues VST

Issue / Primary	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance	Reviewed Non Conformity /	Determined Cause of Non	Evaluated action to	Planning and documenting	Verify Action does not	Effectiveness of corrective
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ID / Call ID		or Complaint / No preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / or dealt with in s	Complaint and determine if its a vigilance Issue requiring a corrective action plan	Conformity / Complaint	Ensure does not recur	action needed and implementation QC 28b	adversely affect Safety Performance or regulatory requirements	action reviewed
<a href="#">335085</a> <a href="#">12 Jul 2024</a>	Shipped Items Return to Supplier BOX988	<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 BOX988 warrant a NON conformance report via the CAPA process VM3COP10

**12 Jul 2024 Derek Lamb**

vst normal sensor fail types

Non Conformance Issues

VST

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 / or dealt with in s	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="#">334360</a> <a href="#">05 Jul 2024</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

**15 Jul 2024 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb Checked back through all Non Conformance issues (not automatically generated) to 1st June 24. Nothing of concern or to be investigated or reported on. 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 issue, nothing to worry about. Non Conformance, complaints and feedback headers reviewed 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

**16 Jul 2024 Derek Lamb**

thankyou

Audit Training

VST

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 / or dealt with in s	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="#">333816</a> <a href="#">01 Jul 2024</a>	Audit 08 Training VST (184)	<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 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

**05 Jul 2024 Derek Lamb**

Next Action Changed From Derek Lamb To Michael Lamb

**19 Sep 2024 Michael Lamb**

Next Action Changed From Michael Lamb To Derek Lamb Training audit completed and attached. Nothing Outstanding Please review

**19 Sep 2024 Derek Lamb**

thankyou

Audit Repairs and Service

VST

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 / or dealt with in s	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="#">333815</a> <a href="#">01 Jul 2024</a>	Audit 11 Repairs And Service VST (179)	<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

**06 Aug 2024 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb completed audit attached no outstanding issue. Please review

**14 Aug 2024 Derek Lamb**  
DoneNon Conformance Issues  
VST

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 / or dealt with in s	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="#">333619</a> <a href="#">27 Jun 2024</a>	Shipped Items Return to Supplier BOX985	<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 BOX985 warrant a NON conformance report via the CAPA process VM3COP10

**01 Jul 2024 Derek Lamb**  
vst o2 sensors normal fail typeAudit Documentation Control  
VST

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 / or dealt with in s	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="#">331857</a> <a href="#">10 Jun 2024</a>	Audit 10 Documentation Control VST (183)	<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 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

**10 Jul 2024 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb completed audit attached no outstanding issues

**12 Jul 2024 Derek Lamb**  
thankyouNon Conformance Issues  
VST

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	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	Effectiveness of corrective action reviewed
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		Requested / Product Failure but no requirement to Escalate Non conformance / or dealt with in s	corrective action plan				regulatory requirements	
<a href="#">331849</a> <a href="#">07 Jun 2024</a>	UnShipped Items Return to Supplier BOX977	<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 BOX977 warrant a NON conformance report via the CAPA process VM3COP10

**17 Jun 2024 Derek Lamb**

vst sensor no cap required

## Non Conformance Issues

VST

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 / or dealt with in s	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="#">331493</a> <a href="#">05 Jun 2024</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

**10 Jun 2024 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb Checked back through all Non Conformance issues (not automatically generated) to 1st May 24. Nothing of concern or to be investigated or reported on. 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 issue, nothing to worry about. Non Conformance, complaints and feedback headers reviewed 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

**14 Jun 2024 Derek Lamb**

thankyou

## Non Conformance Issues

VST

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 / or dealt with in s	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="#">331399</a> <a href="#">04 Jun 2024</a>	Shipped Items Return to Supplier BOX978	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Steve Nixon

Does this Return BOX978 warrant a NON conformance report via the CAPA process VM3COP10

**04 Jun 2024 Derek Lamb**

Next Action Changed From Derek Lamb To Steve Nixon

Priority Changed From 5 To 1 i dont see a fault code of anything with these, are you aware or the return

## Non Conformance Issues

VST

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 / or dealt with in s	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="#">330478</a> <a href="#">23 May</a>	Shipped Items Return to Supplier BOX976	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

2024								
Derek Lamb Does this Return BOX976 warrant a NON conformance report via the CAPA process VM3COP10								
24 May 2024 Derek Lamb vst sensor normal fail type no cap required								
Audit Calibration VST								
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 / or dealt with in s	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="#">329966</a> <a href="#">20 May 2024</a>	Audit 06 Calibration VST (182)	<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 06 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								
03 Jun 2024 Helen Lamb Next Action Changed From Helen Lamb To Michael Lamb current audit attached with up to date processes we use the same doc for viamed and VST								
12 Jun 2024 Helen Lamb Next Action Changed From Michael Lamb To Derek Lamb completed audit attached, carried out by myself and Michael No issues or anything outstanding								
12 Jul 2024 Derek Lamb thankyou								
Non Conformance Issues VST								
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 / or dealt with in s	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="#">328650</a> <a href="#">07 May 2024</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								
10 May 2024 Helen Lamb Next Action Changed From Helen Lamb To Derek Lamb Checked back through all Non Conformance issues (not automatically generated) to 1st April 24. Nothing of concern or to be investigated or reported on. 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. Non Conformance issues review screen - No issues. No other feedback issues relating to non conformances that need to be monitored or reviewed. Order Invoice Error Logs - no issue, nothing to worry about. Derek please review the above								
16 May 2024 Derek Lamb thankyou								
Non Conformance Issues VST								
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 / or dealt with in s	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="#">328341</a> <a href="#">03 May 2024</a>		<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
General 03 May 2024 Derek Lamb just internal normal nonconformance reviews								
Complaints from VST to a Supplier VST								
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 / or dealt with in s	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="#">328338</a> <a href="#">03 May 2024</a>		<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
General 03 May 2024 Derek Lamb no complaints								
Complaints VST								
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 / or dealt with in s	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="#">328336</a> <a href="#">03 May 2024</a>		<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
General 03 May 2024 Derek Lamb no complaints								
Audit Production VST								
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 / or dealt with in s	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="#">328012</a> <a href="#">01 May 2024</a>	Audit 15 Production VST (175)	<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 15 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								
<b>07 May 2024 Helen Lamb</b>								
Next Action Changed From Helen Lamb To Derek Lamb completed audit attached no issues. Please review. If you want ML to review this please pass on and let him know.								
<b>08 May 2024 Derek Lamb</b> thankyou								
Audit Post Marketing Surveillance VST								
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	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	Effectiveness of corrective action reviewed

		Requested / Product Failure but no requirement to Escalate Non conformance / or dealt with in s	corrective action plan				regulatory requirements	
<a href="#">327858</a> <a href="#">30 Apr 2024</a> 329104	Audit 22 Post Market Surveillance VST (180)	<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 22 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 2024 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb completed audit attached, issues outstanding 327860, 327234, 311115 all referenced in issue 329104 sent relating to the Viamed Audit Please review

**16 Oct 2024 Derek Lamb**

all closed now thankyou

Non Conformance Issues

VST

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 / or dealt with in s	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="#">327683</a> <a href="#">26 Apr 2024</a>	Shipped Items Return to Supplier BOX974	<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 BOX974 warrant a NON conformance report via the CAPA process VM3COP10

**26 Apr 2024 Derek Lamb**

vst sensors , normal fail types

Non Conformance Issues

VST

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 / or dealt with in s	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="#">327682</a> <a href="#">26 Apr 2024</a>	Shipped Items Return to Supplier BOX971	<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 BOX971 warrant a NON conformance report via the CAPA process VM3COP10

**26 Apr 2024 Derek Lamb**

vst sensor high output, no cap required

Audit Handling and Storage

VST

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 / or dealt with in s	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="#">326414</a> <a href="#">15 Apr 2024</a>	Audit 07 Handling And Storage VST (178)	<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 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

**26 Apr 2024 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb Audit completed and attached, no issues outstanding please review

**06 Jun 2024 Derek Lamb**

thankyou

Non Conformance Issues  
VST

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 / or dealt with in s	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="#">325635</a> <a href="#">05 Apr 2024</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

**09 Apr 2024 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb VST Checked back through all Non Conformance issues (not automatically generated) to 1st March 24. Nothing of concern or to be investigated or reported on. 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. Non Conformance issues review screen - No issues. No other feedback issues relating to non conformances that need to be monitored or reviewed. Non conformance meeting header - Several positive feedbacks have been added. No issues or areas of concern. Order Invoice Error Logs - no issue, nothing to worry about. Derek please review the above

**10 Apr 2024 Derek Lamb**

thankyou

Audit CE Files  
VST

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 / or dealt with in s	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="#">324483</a> <a href="#">25 Mar 2024</a>	Audit 12 CE Files VST (176)	<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 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

**16 May 2024 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb completed audit attached, no issues. Derek please review

**16 May 2024 Derek Lamb**

thankyou

Non Conformance Issues  
VST

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 / or dealt with in s	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="#">324145</a> <a href="#">20 Mar 2024</a>	Shipped Items Return to Supplier BOX967	<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 BOX967 warrant a NON conformance report via the CAPA process VM3COP10								
<b>26 Mar 2024 Derek Lamb</b> normal vst sensor return no capa required								
Audit Goods Inwards and Product Identity VST								
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 / or dealt with in s	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="#">323163</a> <a href="#">11 Mar 2024</a>	Audit 09 Goods Inward And Product Identity VST (174)	<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								
<b>26 Apr 2024 Helen Lamb</b> Next Action Changed From Helen Lamb To Derek Lamb Completed audit attached Issue 327056 sent re process 673 and 674. Nothing else outstanding. Please review								
<b>17 May 2024 Derek Lamb</b> thankyou								
Non Conformance Issues VST								
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 / or dealt with in s	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="#">322561</a> <a href="#">05 Mar 2024</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								
<b>11 Mar 2024 Helen Lamb</b> Next Action Changed From Helen Lamb To Derek Lamb Checked back through all Non Conformance issues (not automatically generated) to 1st Feb 24. Nothing of concern or to be investigated or reported on. 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. Non Conformance issues review screen - No issues. No other feedback issues relating to non conformances that need to be monitored or reviewed. Non conformance meeting header - Several positive feedbacks have been added. No issues or areas of concern. Order Invoice Error Logs - no issue, nothing to worry about. Derek please review the above								
<b>13 Mar 2024 Derek Lamb</b> thankyou								
Non Conformance Issues VST								
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 / or dealt with in s	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="#">319657</a> <a href="#">05 Feb</a> <a href="#">2024</a>	Non conformance review history VST (286)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
<p>Derek Lamb System Generated Check the below review is being carried out</p> <p>Check the history of the last Non conformance review,</p> <p>check actions are being carried out, and non conformances are not reoccurring</p> <p><b>05 Feb 2024 Derek Lamb</b> vst non conformances tasks upto date, no major issues found</p>								
Non Conformance Issues VST								
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 / or dealt with in s	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="#">319656</a> <a href="#">05 Feb</a> <a href="#">2024</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"/>
<p>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</p> <p><b>05 Feb 2024 Helen Lamb</b> Checked back through all Non Conformance issues (not automatically generated) to 1st Jan 23. Nothing of concern or to be investigated or reported on. 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. Non Conformance issues review screen - No issues. No other feedback issues relating to non conformances that need to be monitored or reviewed. Non conformance meeting header - Update of an objective in the management review, no other issues or areas of concern. Order Invoice Error Logs - no issue, nothing to worry about. Derek please review the above</p>								
Adverse incident reporting VST								
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 / or dealt with in s	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="#">319554</a> <a href="#">02 Feb</a> <a href="#">2024</a>		<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
<p>General 02 Feb 2024 Derek Lamb no incidents</p>								
Audits Closure VST								
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 / or dealt with in s	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="#">318763</a> <a href="#">25 Jan</a> <a href="#">2024</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"/>
<p>General 25 Jan 2024 Derek Lamb closed lat years audits, agreed to next year 2024 schedula</p>								

Results of internal audits / Mini Audits VST								
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 / or dealt with in s	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="#">318761</a> <a href="#">25 Jan 2024</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  
25 Jan 2024 Derek Lamb  
happy with last years audits, and agreed to next yers schedule

Non Conformance Issues VST								
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 / or dealt with in s	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="#">318651</a> <a href="#">25 Jan 2024</a>	Shipped Items Return to Supplier BOX960	<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 BOX960 warrant a NON conformance report via the CAPA process VM3COP10

**26 Jan 2024 Derek Lamb**  
vst normal sensor fail type no capa required

Non Conformance Issues VST								
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 / or dealt with in s	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="#">316859</a> <a href="#">05 Jan 2024</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

**09 Jan 2024 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb Checked back through all Non Conformance issues (not automatically generated) to 1st Jan 23. Nothing of concern or to be investigated or reported on. 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. Non Conformance issues review screen - No issues. No other feedback issues relating to non conformances that need to be monitored or reviewed. Non conformance meeting header - Nothing in last month. Nothing further. Order Invoice Error Logs - no issue, nothing to worry about. Derek please review the above

**09 Jan 2024 Derek Lamb**  
Done

Audit Picking and Packing VST								
Issue / Primary	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No	Reviewed Non Conformity / Complaint and	Determined Cause of Non	Evaluated action to Ensure	Planning and documenting action needed and	Verify Action does not adversely affect	Effectiveness of corrective

ID / Call ID		preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / or dealt with in s	determine if its a vigilance Issue requiring a corrective action plan	Conformity / Complaint	does not recur	implementation QC 28b	Safety Performance or regulatory requirements	action reviewed
<a href="#">316318</a> <a href="#">01 Jan 2024</a>	Audit 01 Picking Packing VST (194)	<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 01 due 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

**08 Jan 2024 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb Completed audit attached no issues please review

**09 Jan 2024 Derek Lamb**  
thankyouAudit Design Control  
VST

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 / or dealt with in s	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="#">316317</a> <a href="#">01 Jan 2024</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"/>

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

**05 Feb 2024 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb audit attached no ongoing issues please review

**06 Feb 2024 Derek Lamb**  
thasnkyouAudit Internal Audits  
VST

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 / or dealt with in s	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="#">316128</a> <a href="#">28 Dec 2023</a> 313880	Audit 17 Internal Audits VST (191) Task 1094 not completed	<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 Helen Lamb sent to Derek Lamb

Audit 17 Internal Audits VST (191) Task 1094 not completed Board meeting has been delayed as a Director has been really unwell. Will be carried out in January 24

**31 Jan 2024 Derek Lamb**  
has been done now,Non Conformance Issues  
VST

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 / or dealt with in s						
<a href="#">314196</a> <a href="#">05 Dec 2023</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"/>
<p>Derek Lamb</p> <p>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</p>								
<p><b>14 Dec 2023 Helen Lamb</b></p> <p>Next Action Changed From Helen Lamb To Derek Lamb Checked back through all Non Conformance issues (not automatically generated) to 1st Dec 23. Nothing of concern or to be investigated or reported on. 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. Non Conformance issues review screen - No issues. No other feedback issues relating to non conformances that need to be monitored or reviewed. Non conformance meeting header - Nothing in last month. Nothing further. Order Invoice Error Logs - no issue, nothing to worry about. Derek please review the above</p>								
<p><b>15 Dec 2023 Derek Lamb</b></p> <p>thankyou</p>								
<p>Audit of Audits VST</p>								
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 / or dealt with in s	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="#">313881</a> <a href="#">01 Dec 2023</a>	Audit 21 Audit Of Audit VST (192)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
<p>Derek Lamb</p> <p>System Generated Audit 21 due Review the Audit Calendar Screen ISO -&gt; Audit Calendar Complete Audit 21 Confirm if Audit calendar needs changing. 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</p>								
<p><b>29 Dec 2023 Derek Lamb</b></p> <p>Next Action Changed From Derek Lamb To Helen Lamb ive given you the audit to scan and attach to this issue please</p>								
<p><b>02 Jan 2024 Helen Lamb</b></p> <p>Next Action Changed From Helen Lamb To Derek Lamb Completed audit attached, completed by you. no issues please check correct is attached.</p>								
<p><b>02 Jan 2024 Derek Lamb</b></p>								
<p>Audit Internal Audits VST</p>								
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 / or dealt with in s	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="#">313880</a> <a href="#">01 Dec 2023</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"/>
<p>Derek Lamb</p> <p>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. 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</p>								
<p><b>28 Dec 2023 Helen Lamb</b></p> <p>Created Related Issue #316128 Added by Helen Lamb sent to Derek Lamb Audit 17 Internal Audits VST (191) Task 1094 not completed Board meeting has been delayed as a Director has been really unwell. Will be carried out in January 24</p>								
<p><b>28 Dec 2023 Helen Lamb</b></p> <p>Next Action Changed From Helen Lamb To Derek Lamb Audit completed issue 316128 sent re task not completed. Please review the attached.</p>								



Non Conformance Issues VST								
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 / or dealt with in s	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="#">313647</a> <a href="#">28 Nov 2023</a>	Shipped Items Return to Supplier BOX941	<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 BOX941 warrant a NON conformance report via the CAPA process VM3COP10

**28 Nov 2023 Derek Lamb**

80100082240983M01112018No Output vst normal sensor type fail no capa required

Non Conformance Issues VST								
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 / or dealt with in s	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="#">311635</a> <a href="#">06 Nov 2023</a> 311426	Non-Conformance Vandgraph Website	<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 Derek Lamb sent to Helen Lamb

See Main Issue

**09 Nov 2023 Helen Lamb**

have spoken to Mike and advised him these need to be carried out more regularly

Non Conformance Issues VST								
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 / or dealt with in s	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="#">311536</a> <a href="#">06 Nov 2023</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

**15 Nov 2023 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb Checked back through all Non Conformance issues (not automatically generated) to 1st Nov 23. Nothing of concern or to be investigated or reported on. 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. Non Conformance issues review screen - No issues. No other feedback issues relating to non conformances that need to be monitored or reviewed. Non conformance meeting header - Nothing in last month. Nothing further. Order Invoice Error Logs - no issue, nothing to worry about. Derek please review the above

**16 Nov 2023 Derek Lamb**

Done

Non Conformance Issues VST								
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	Determined Cause of Non Conformity / Complaint	Evaluated action to Ensure	Planning and documenting action needed and	Verify Action does not adversely affect Safety	Effectiveness of corrective action reviewed

		actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / or dealt with in s	vigilance Issue requiring a corrective action plan		does not recur	implementation QC 28b	Performance or regulatory requirements	
<a href="#">311426</a> <a href="#">03 Nov 2023</a>	Non-Conformance Vandagraph Website	<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 Derek Lamb

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

Order have not been marked as fulfilled from 18th October for 13 Vandagraph orders. I will go through them now to check system and fulfil.

**06 Nov 2023 Derek Lamb**

Created Related Issue #311635

Added by Derek Lamb sent to Helen Lamb

See Main Issue

## Audit Organisation and Process Verification

VST

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 / or dealt with in s	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="#">311116</a> <a href="#">01 Nov 2023</a>	Audit 20 Process Verification To Managment VST (181)	<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 20 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

**05 Dec 2023 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb Completed audit attached no outstanding issues. Please review

**05 Dec 2023 Derek Lamb**

thankyou

## Audit Management Review

VST

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 / or dealt with in s	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="#">309399</a> <a href="#">16 Oct 2023</a>	Audit 18 Management Review VST (188)	<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 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. 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

**15 Nov 2023 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb Completed Audit Attached please review. Assessed from the 2022 Management review, 2023 is now due and planned for next month. Nothing outstanding

**15 Nov 2023 Helen Lamb**

i labelled the attached doc wrong its for 2023

**16 Nov 2023 Derek Lamb**

Done

**16 Nov 2023 Derek Lamb**

Done

Complaints from VST to a Supplier

VST

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 / or dealt with in s	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="#">309020</a> <a href="#">10 Oct 2023</a>	test	<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  
test

**11 Oct 2023 Derek Lamb**  
Done

#### Future Reviews - Internal Audits VST

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 / or dealt with in s	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="#">308987</a> <a href="#">10 Oct 2023</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  
10 Oct 2023 Derek Lamb  
on going

#### VST Customer Complaints VST

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 / or dealt with in s	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="#">308961</a> <a href="#">10 Oct 2023</a>	test	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>

Derek Lamb  
10 Oct 2023 Derek Lamb  
test issue

**11 Oct 2023 Derek Lamb**  
Done

#### Non Conformance Issues VST

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 / or dealt with in s	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="#">308534</a> <a href="#">05 Oct 2023</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"/>

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

**05 Oct 2023 Helen Lamb**

Please see issue 305778. September issue was done late due to illness, so covers Octobers Non conformance review.

**05 Oct 2023 Helen Lamb**

Please see issue 305778. September issue was done late due to illness, so covers Octobers Non conformance review.

## Non Conformance Issues

VST

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 / or dealt with in s	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="#">307437</a> <a href="#">22 Sep 2023</a>	Shipped Items Return to Supplier BOX938	<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 BOX938 warrant a NON conformance report via the CAPA process VM3COP10

**27 Sep 2023 Derek Lamb**

vst o2 sensor unstable output, SN aware, normal fail type, no cap required at this time

## Non Conformance Issues

VST

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 / or dealt with in s	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="#">305816</a> <a href="#">05 Sep 2023</a>	Shipped Items Return to Supplier BOX930	<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 BOX930 warrant a NON conformance report via the CAPA process VM3COP10

**05 Sep 2023 Derek Lamb**

VST normal sensor fail type Done

## Non Conformance Issues

VST

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 / or dealt with in s	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="#">305778</a> <a href="#">05 Sep 2023</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

**03 Oct 2023 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb Checked back through all Non Conformance issues (not automatically generated) to 1st Oct 23. I have had Covid so this has been delayed this will cover the next issue too. So August and September 23. Nothing of concern or to be investigated or reported on. 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. Non Conformance issues review screen - No issues. No other feedback issues relating to non conformances that need to be monitored or reviewed. Non conformance meeting header - Nothing in last two months. Nothing further. Order Invoice Error Logs - no issue, nothing to worry about. Derek please review the above

**04 Oct 2023 Derek Lamb**

thankyou

Audit Purchasing  
VST

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 / or dealt with in s	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="#">305390</a> <a href="#">01 Sep 2023</a>	Audit 05 Purchasing Suppliers VST (190)	<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

**12 Sep 2023 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb Completed audit attached no issues please review

**15 Sep 2023 Derek Lamb**

thankyou

Audit Contract Review  
VST

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 / or dealt with in s	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="#">305388</a> <a href="#">01 Sep 2023</a>	Audit 02 Contract Review VST (187)	<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

**11 Sep 2023 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb Audit completed and no issues please review the attached.

**11 Sep 2023 Derek Lamb**

thankyou

Non Conformance Issues  
VST

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 / or dealt with in s						
<a href="#">303194</a> <a href="#">07 Aug 2023</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"/>
<p>Derek Lamb</p> <p>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</p>								
<p><b>07 Aug 2023 Helen Lamb</b></p> <p>Next Action Changed From Helen Lamb To Derek Lamb Checked back through all Non Conformance issues (not automatically generated) to 1st July 23. Nothing of concern or to be investigated or reported on. 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. Non Conformance issues review screen - No issues. No other feedback issues relating to non conformances that need to be monitored or reviewed. Non conformance meeting header - 300731 issue wrong invoice sent to customer, human error, relevant staff have been spoken to. Nothing further. Order Invoice Error Logs - no issue, nothing to worry about. Derek please review the above</p>								
<p><b>07 Aug 2023 Derek Lamb</b></p> <p>thankyou</p>								
<p>Audit Health and Saftey VST</p>								
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 / or dealt with in s	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="#">302436</a> <a href="#">01 Aug 2023</a>	Audit 19 Health And Saftey VST (186)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
<p>Derek Lamb</p> <p>System Generated Do HSE Audit Audit No 19 Review Last years Audit see if its still suitable 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</p>								
<p><b>01 Aug 2023 Helen Lamb</b></p> <p>Next Action Changed From Helen Lamb To Derek Lamb Completed audit attached, please review. No outstanding issues.</p>								
<p><b>02 Aug 2023 Derek Lamb</b></p> <p>Thankyou Done</p>								
<p>Non Conformance Issues VST</p>								
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 / or dealt with in s	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="#">301436</a> <a href="#">19 Jul 2023</a> 299479	Audit 11 Repairs And Service VST (179) Non Conformance QA records	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
<p>Derek Lamb</p> <p>Added by Helen Lamb sent to Derek Lamb</p> <p>Test records have QA but not in the QA Data base. system to be updated</p>								
<p><b>21 Jul 2023 Derek Lamb</b></p> <p>Linked TaskID Changed From 0 To 456</p>								
<p><b>18 Sep 2023 Derek Lamb</b></p> <p>updated the stock references to havea pass fail and output gielfs to appear in the SRS system.</p>								
<p>Audit Analysis of Data VST</p>								

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 / or dealt with in s	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="#">300839</a> <a href="#">13 Jul 2023</a>	Audit 23 Analysis Of Data VST (185)	<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 23 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 Jul 2023 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb Completed audit attached. Please check, the only issue outstanding is the one for the management review which cannot be completed until signing off in the next meeting.

**31 Jul 2023 Helen Lamb**

**31 Jul 2023 Derek Lamb**  
thankyou

Non Conformance Issues  
VST

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 / or dealt with in s	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="#">300820</a> <a href="#">12 Jul 2023</a> 300732	Order Error : 143286 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

Added by Helen Lamb sent to Helen Lamb  
Incorrect invoice sent to customer, sent invoice for M3S RST142896-1 instead of RST143286-1. We have sent the wrong invoice to a VST customer. Please be extra careful when adding invoices to your emails. Especially VST ones.

**08 Aug 2023 Helen Lamb****08 Aug 2023 Helen Lamb**

Non Conformance Issues  
VST

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 / or dealt with in s	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="#">300732</a> <a href="#">12 Jul 2023</a>	Order Error : 143286 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 143286

Order Entered by Janine Gill

Order Checked by Kate Griffiths

Office

Error was New Error

Fault:

Incorrect invoice sent to customer, sent invoice for M3S RST142896-1 instead of RST143286-1. Non-Conformance raised issue #300731

Possible Fix

**12 Jul 2023 Helen Lamb**

Created Related Issue #300820

Added by Helen Lamb sent to Helen Lamb

Incorrect invoice sent to customer, sent invoice for M3S RST142896-1 instead of RST143286-1. We have sent the wrong invoice to a VST customer. Please be extra careful when adding invoices to your emails. Especially VST ones.

#### Non Conformance Issues VST

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 / or dealt with in s	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="#">300731</a> <a href="#">12 Jul 2023</a>	rEvo x M3S Incorrect Invoice Sent to Customer	<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 Derek Lamb

Incorrect invoice sent to customer, sent invoice for M3S RST142896-1 instead of RST143286-1 for rEvo. Also raised as Log Error against rEvo order.

**07 Aug 2023 Derek Lamb**

first ever occurrence of this - looks like human error, spoken to office staff no further action taken

#### Non Conformance Issues VST

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 / or dealt with in s	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="#">299811</a> <a href="#">05 Jul 2023</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

**10 Jul 2023 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb Checked back through all Non Conformance issues (not automatically generated) to 1st June 23. Nothing of concern or to be investigated or reported on. 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. No other feedback issues relating to non conformances that need to be monitored or reviewed. Non conformance meeting header - nothing new Order Invoice Error Logs - no issue, nothing to worry about. Derek please review the above

**11 Jul 2023 Derek Lamb**

Done

#### Audit Repairs and Service VST

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 / or dealt with in s	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="#">299479</a> <a href="#">03 Jul 2023</a>	Audit 11 Repairs And Service VST (179)	<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

**19 Jul 2023 Helen Lamb**

Created Related Issue #301436

Added by Helen Lamb sent to Derek Lamb

Test records have QA but not in the QA Data base. system to be updated

**10 Aug 2023 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb completed audit attached, no issues

**25 Aug 2023 Derek Lamb**

thankyou

Audit Training  
VST

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 / or dealt with in s	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="#">299315</a> <a href="#">30 Jun 2023</a>	Audit 08 Training VST (184)	<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 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 2023 Derek Lamb**

Next Action Changed From Derek Lamb To Helen Lamb

**24 Jul 2023 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb completed audit attached that you carried out. Please check it scanned ok

**24 Jul 2023 Derek Lamb**

Done

Non Conformance Issues  
VST

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 / or dealt with in s	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="#">299310</a> <a href="#">29 Jun 2023</a>	Shipped Items Return to Supplier BOX921	<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 BOX921 warrant a NON conformance report via the CAPA process VM3COP10

**29 Jun 2023 Derek Lamb**

reviewed vst , normal fail types

**29 Jun 2023 Derek Lamb**

reviewed vst , normal fail types

Audit Documentation Control  
VST

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 / or dealt with in s	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="#">297581</a> <a href="#">12 Jun 2023</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

**26 Jun 2023 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb audit done no issues

**26 Jun 2023 Derek Lamb**

thankyou

Non Conformance Issues

VST

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 / or dealt with in s	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="#">296962</a> <a href="#">05 Jun 2023</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

**09 Jun 2023 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb Checked back through all Non Conformance issues (not automatically generated) to 1st May 23. Issue 295394 QC21 forms generated for BSI minors, not a problem and nothing else of concern or to be investigated or reported on. 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. No other feedback issues relating to non conformances that need to be monitored or reviewed. Order Invoice Error Logs - no issue, nothing to worry about. Derek please review the above

**13 Jun 2023 Derek Lamb**

Done

VST Customer Complaints

VST

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 / or dealt with in s	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="#">296752</a> <a href="#">01 Jun 2023</a>	Divesoft - complaint list number 23OB0251	<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 Steve Hardaker

Concerns 8010020 oxygen sensors Please process through the updated complaints system. See the attached emails. I will get back to Divesoft with a summary of the findings concerning the overspill of the conformal coating and also process the credits relating to the SRS.

**02 Jun 2023 Steve Nixon**

Credit done TST143790-0

**02 Jun 2023 Steve Hardaker**

There are no attached emails, please can you upload them and bounce this issue back to me.

**22 Jun 2023 Derek Lamb**

Linked TaskID Changed From 0 To 1193

**14 Aug 2023 Steve Hardaker**

Customer Complain Report CCR154 was been generated. Just need to print along with supporting documents and add to the paper file.

**16 Aug 2023 Steve Hardaker**

Next Action Changed From Steve Hardaker To Derek Lamb Added to paper file in office meeting room. I believe this is now complete, just requires signing off by a Director.

**23 Aug 2023 Derek Lamb**

Done

Audit Calibration

VST

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	Determined Cause of Non Conformity / Complaint	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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		actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / or dealt with in s	vigilance Issue requiring a corrective action plan		does not recur	implementation QC 28b	Performance or regulatory requirements	
<a href="#">295678</a> <a href="#">22 May 2023</a>	Audit 06 Calibration VST (182)	<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 06 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

**23 Jun 2023 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb completed Audit attached no issues

**26 Jun 2023 Derek Lamb**

thankyou

Non Conformities Review  
VST

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 / or dealt with in s	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="#">295327</a> <a href="#">16 May 2023</a>	2342435-202305-N2 ISO 9001:2015 9.2 Minor	<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"/>

Helen Lamb

Added by Derek Lamb sent to Derek Lamb

1. The unique identifier (from the BSI report and also any internal reference). 2. The statement of Nonconformity as written in the BSI report. 3. Root Cause Analysis. 4. Relevant Immediate Correction (where applicable). 5. Relevant and Proportionate Corrective Action. 6. Person responsible to complete the action(s). 7. Time for completion of all identified actions. The process for internal audit is not fully effective as it was not clear that all issues raised during internal audit were part of the non- conformity review process as per documented procedure 9.2 Internal audit Picking packing audit 2023

**16 May 2023 Derek Lamb**

Next Action Changed From Derek Lamb To Helen Lamb

**18 May 2023 Helen Lamb**

HL revision added 295327.12614 qc 21 initial document N2\_18\_05\_23 HL1

**21 May 2024 Helen Lamb**

Effectiveness has been reviewed and completed. QC21 Non conformance has been closed by Derek Lamb today.

**21 Jun 2024 Helen Lamb**Non Conformities Review  
VST

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 / or dealt with in s	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="#">295321</a> <a href="#">16 May 2023</a>	2342435-202305-N1 ISO 9001:2015 5.2 Minor	<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"/>

Helen Lamb

Added by Derek Lamb sent to Derek Lamb

1. The unique identifier (from the BSI report and also any internal reference). 2. The statement of Nonconformity as written in the BSI report. 3. Root Cause Analysis. 4. Relevant Immediate Correction (where applicable). 5. Relevant and Proportionate Corrective Action. 6. Person responsible to complete the action(s). 7. Time for completion of all identified actions. The process for the quality policy is not fully effective as it was defined as a secondary level document. 22062 VM3COP.00.00 Company quality policy 16 sept 2017

**16 May 2023 Derek Lamb**

Next Action Changed From Derek Lamb To Helen Lamb

**18 May 2023 Helen Lamb**

section 5.2 of 9001 first revision by me doc named 295321.12613 qc21 initial document N1\_18\_05\_23 HL1

**21 May 2024 Helen Lamb**

effectiveness verified and closed by Derek Lamb doc attached.

**21 Jun 2024 Helen Lamb**Non Conformance Issues  
VST

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 / or dealt with in s	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="#">295019</a> <a href="#">12 May 2023</a>	Shipped Items Return to Supplier BOX915	<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 BOX915 warrant a NON conformance report via the CAPA process VM3COP10

**15 May 2023 Derek Lamb**

normal sensor type fails

Non Conformance Issues  
VST

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 / or dealt with in s	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="#">294268</a> <a href="#">05 May 2023</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

**15 May 2023 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb Checked back through all Non Conformance issues (not automatically generated) to 1st April 23 nothing of concern or to be investigated or reported on. BSI were in last week for 9001:2015 and four Minors were given, these will be investigated and processed shortly. 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. No other feedback issues relating to non conformances that need to be monitored or reviewed. Order Invoice Error Logs - no issue, nothing to worry about. Derek please review the above

**15 May 2023 Derek Lamb**

thankyou

Audit Post Marketing Surveillance  
VST

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 / or dealt with in s	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="#">293724</a> <a href="#">02 May 2023</a>	Audit 22 Post Market Surveillance VST (180)	<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 22 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

**02 Jun 2023 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb completed audit attached, no issues from this VST audit

<b>05 Jun 2023 Derek Lamb</b> thankouy								
Audit Production VST								
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 / or dealt with in s	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="#">293720</a> <a href="#">02 May 2023</a>	Audit 15 Production VST (175)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
<b>Derek Lamb</b> System Generated Audit 15 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								
<b>02 Jun 2023 Helen Lamb</b>								
Next Action Changed From Helen Lamb To Derek Lamb completed audit attached , no issues and no production carried out by VST								
<b>02 Jun 2023 Derek Lamb</b> thankyou								
Non Conformance Issues VST								
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 / or dealt with in s	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="#">292702</a> <a href="#">18 Apr 2023</a>	Shipped Items Return to Supplier BOX912	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
<b>Derek Lamb</b> Does this Return BOX912 warrant a NON conformance report via the CAPA process VM3COP10								
<b>28 Apr 2023 Derek Lamb</b> Done								
Audit Handling and Storage VST								
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 / or dealt with in s	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="#">292443</a> <a href="#">17 Apr 2023</a>	Audit 07 Handling And Storage VST (178)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
<b>Derek Lamb</b> System Generated Audit 07 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								
<b>19 Jun 2023 Helen Lamb</b>								
Next Action Changed From Helen Lamb To Derek Lamb audit completed no issues please review Done								
<b>19 Jun 2023 Derek Lamb</b> thankyou								
Non Conformance Issues								

VST								
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 / or dealt with in s	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="#">291403</a> <a href="#">05 Apr 2023</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

**03 May 2023 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb Checked back through all Non Conformance issues (not automatically generated) to 1st March 23 nothing of concern or to be investigated or reported on. 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. No other feedback issues relating to non conformances that need to be monitored or reviewed. Order Invoice Error Logs - no issue, nothing to worry about. Derek please review the above

**09 May 2023 Derek Lamb**

thankyou

## Audit CE Files

VST								
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 / or dealt with in s	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="#">290136</a> <a href="#">22 Mar 2023</a>	Audit 12 CE Files VST (176)	<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 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

**29 Mar 2023 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb Scanned to issue completed by Steve Nixon and Derek Lamb. Attached.

**30 Mar 2023 Derek Lamb**

thankyou

**14 Sep 2023 Derek Lamb**

## Audit Goods Inwards and Product Identity

VST								
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 / or dealt with in s	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="#">288861</a> <a href="#">09 Mar 2023</a>	Audit 09 Goods Inward And Product Identity VST (174)	<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

**13 Apr 2023 Helen Lamb**

Created Related Issue #292109

Added by Helen Lamb sent to Derek Lamb

there are two SRSs without email addresses All the other info is there but i though this should probably be in here too. I am unsure if this is a system thing not pulling it in or a person thing

**13 Apr 2023 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb completed audit attached one issue 292109 re SRS info.

**18 Apr 2023 Derek Lamb**

thankyou

Non Conformance Issues

VST

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 / or dealt with in s	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="#">288494</a> <a href="#">06 Mar 2023</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

**22 Mar 2023 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb Checked back through all Non Conformance issues (not automatically generated) to 1st February 23 nothing of concern or to be investigated or reported on. 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. No other feedback issues relating to non conformances that need to be monitored or reviewed. Order Invoice Error Logs - no issue, nothing to worry about. Derek please review the above

**23 Mar 2023 Derek Lamb**

thankyou

Non Conformance Issues

VST

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 / or dealt with in s	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="#">284669</a> <a href="#">25 Jan 2023</a> 246776	Audit 01 Picking Packing VST (194) Needs a re write as system been updated	<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

Both Viamed and VSt need a re write as reflects old system

**17 Feb 2023 Helen Lamb**

Priority Changed From 5 To 1

**11 May 2023 Derek Lamb**

Header Changed From 614 VST Audits Calander Audit Picking and Packing To VST Management Non Conformance Issues

**26 Jun 2023 Helen Lamb**

Done

**26 Jun 2023 Helen Lamb**

Audit Picking and Packing

VST

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	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	Effectiveness of corrective action reviewed
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		Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / or dealt with in s	requiring a corrective action plan				regulatory requirements	
<a href="#">281965</a> <a href="#">03 Jan</a> <a href="#">2023</a>	Audit 01 Picking Packing VST (194)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
<p>Derek Lamb</p> <p>System Generated Audit 01 due 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</p>								
<p><b>16 Feb 2023 Helen Lamb</b></p> <p>Next Action Changed From Helen Lamb To Derek Lamb no issues, Audit attached please review</p>								
<p><b>20 Feb 2023 Derek Lamb</b> reviewed</p>								
<p>Audit Design Control VST</p>								
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 / or dealt with in s	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="#">281964</a> <a href="#">03 Jan</a> <a href="#">2023</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"/>
<p>Derek Lamb</p> <p>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</p>								
<p><b>23 Jun 2023 Helen Lamb</b></p> <p>Next Action Changed From Helen Lamb To Derek Lamb you were going to go through this with Steve</p>								
<p><b>20 Sep 2023 Derek Lamb</b></p> <p>Next Action Changed From Derek Lamb To Helen Lamb</p>								
<p><b>21 Sep 2023 Helen Lamb</b></p> <p>Next Action Changed From Helen Lamb To Derek Lamb completed audit attached please can you check its ok . no issues</p>								
<p><b>22 Sep 2023 Derek Lamb</b> THANKYOU</p>								
<p>Audit of Audits VST</p>								
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 / or dealt with in s	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="#">279306</a> <a href="#">01 Dec</a> <a href="#">2022</a>	Audit 21 Audit Of Audit VST (192)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
<p>Helen Lamb</p> <p>System Generated Audit 21 due Review the Audit Calendar Screen ISO -&gt; Audit Calendar Complete Audit 21 Confirm if Audit calendar needs changing. 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</p>								
<p><b>06 Dec 2022 Helen Lamb</b> Scanned Audit attached as per your request.</p>								
<p><b>06 Dec 2022 Derek Lamb</b></p> <p>Next Action Changed From Derek Lamb To Helen Lamb please reveiw and complete</p>								

**06 Dec 2022 Helen Lamb**

Reviewed ok no issues

## Audit Internal Audits

VST

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 / or dealt with in s	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="#">279305</a> <a href="#">01 Dec 2022</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"/>

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. 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

**06 Dec 2022 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb audit completed and reviewed, Was carried out by both of us, no issues

**06 Dec 2022 Derek Lamb**

reviewed thankyou

## Non Conformance Issues

VST

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 / or dealt with in s	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="#">273702</a> <a href="#">05 Oct 2022</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

**28 Oct 2022 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb Checked back through all Non Conformance issues (not automatically generated) to 1st September 22 nothing of concern or to be investigated or reported on. 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. No other feedback issues relating to non conformances that need to be monitored or reviewed. Order Invoice Error Logs - no issue, nothing to worry about. Derek please review the above #276042 issue sent re Customer Complaint and Non Conformance Review Screen as not filtering to VST

**31 Oct 2022 Derek Lamb**

ok, will fix filter on the page

## Non Conformance Issues

VST

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 / or dealt with in s	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="#">270891</a> <a href="#">05 Sep 2022</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

#### 14 Sep 2022 Helen Lamb

Next Action Changed From Helen Lamb To Derek Lamb Checked back through all Non Conformance issues to 1st August 22 nothing of concern or to be investigated or reported on. 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. No other feedback issues relating to non conformances that need to be monitored or reviewed. Order Invoice Error Logs - no issue, nothing to worry about. Derek please review the above

#### 16 Sep 2022 Derek Lamb

Reviewed ok

#### Non Conformance Issues VST

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 / or dealt with in s	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="#">268177</a> <a href="#">05 Aug 2022</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

#### 10 Aug 2022 Helen Lamb

Next Action Changed From Helen Lamb To Derek Lamb Checked back through all Non Conformance issues to 1st July 22 nothing of concern or to be investigated or reported on. 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. No other feedback issues relating to non conformances that need to be monitored or reviewed. Order Invoice Error Logs - no issue, nothing to worry about. Derek please review

#### 10 Aug 2022 Derek Lamb

reviewed

#### Non Conformance Issues VST

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 / or dealt with in s	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="#">267039</a> <a href="#">25 Jul 2022</a> 265294	Non conformance review history VST (285) tasks	<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 Helen Lamb sent to Derek Lamb

Positive feedback re product VST. No Task for this to be reviewed Positive feedback re customer VST. No Task for this to be reviewed Think we should have tasks for these

#### 25 Jul 2022 Helen Lamb

there are two other issues tasks but they seem a general ones. Tasks 1068 and 1069

#### 25 Jul 2022 Derek Lamb

added today as processes with linked tasks

#### 29 Jul 2022 Helen Lamb

#### Non Conformance Issues VST

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 / or dealt with in s						
<a href="#">265294</a> <a href="#">05 Jul</a> <a href="#">2022</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"/>
<p>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</p> <p><b>25 Jul 2022 Helen Lamb</b> Created Related Issue #267039 Added by Helen Lamb sent to Derek Lamb Positive feedback re product VST. No Task for this to be reviewed Positive feedback re customer VST. No Task for this to be reviewed Think we should have tasks for these</p> <p><b>25 Jul 2022 Helen Lamb</b> Checked back through all Non Conformance issues to 1st June 22 nothing of concern or to be investigated or reported on. Review VST Product Feedback Negative (742) nothing new. Positive feedback re product nothing new. No Task for this to be reviewed Review VST Feedback - Customer Feedback Negative (740) nothing new. Positive feedback re customer nothing new. No Task for this to be reviewed Review VST Feedback - Customer Complaints (738) No other feedback issues relating to non conformances that need to be monitored or reviewed. Order Invoice Error Logs - no issue, nothing to worry about.</p>								
Audit Training VST								
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 / or dealt with in s	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="#">264799</a> <a href="#">30 Jun</a> <a href="#">2022</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"/>
<p>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</p> <p><b>14 Jul 2022 Helen Lamb</b> Audit completed Derek, scanned and uploaded by Helen Second scan is in the correct page order</p> <p><b>21 Jul 2022 Derek Lamb</b> Done</p>								
Audit Documentation Control VST								
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 / or dealt with in s	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="#">262667</a> <a href="#">10 Jun</a> <a href="#">2022</a>	Audit 10 Documentation Control VST (183)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
<p>Derek Lamb System Generated 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</p> <p><b>07 Jul 2022 Helen Lamb</b> Next Action Changed From Helen Lamb To Derek Lamb Audit completed no issues, please review</p> <p><b>07 Jul 2022 Derek Lamb</b> thank you</p>								
Non Conformance Issues VST								
Issue / Primary	Subject/Notes	Minor Internal Error i.e. not ISO Non Conformance or Complaint / No	Reviewed Non Conformity / Complaint and	Determined Cause of Non	Evaluated action to Ensure	Planning and documenting action needed and	Verify Action does not adversely affect	Effectiveness of corrective

ID / Call ID		preventative or corrective actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / or dealt with in s	determine if its a vigilance Issue requiring a corrective action plan	Conformity / Complaint	does not recur	implementation QC 28b	Safety Performance or regulatory requirements	action reviewed
<a href="#">262009</a> <a href="#">06 Jun 2022</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"/>

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

**06 Jun 2022 Helen Lamb**

checked back through all Non Conformance issues to 1st May 22 nothing of concern or to be investigated or reported on. Negative feedback re product nothing new. Positive feedback re product nothing new. Negative feedback re customer nothing new. Positive feedback re customer nothing new. No other feedback issues relating to non conformances that need to be monitored or reviewed Order Invoice Error Logs - no issue, nothing to worry about.

Non Conformance Issues  
VST

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 / or dealt with in s	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="#">259203</a> <a href="#">05 May 2022</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"/>

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

**06 May 2022 Helen Lamb**

checked back through all Non Conformance issues to 1st April 22 nothing of concern or to be investigated or reported on. Order Invoice Error Logs - no issue, nothing to worry about. No feedback issues relating to non conformances that need to be monitored or reviewed

Audit Handling and Storage  
VST

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 / or dealt with in s	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="#">257431</a> <a href="#">19 Apr 2022</a>	Audit 07 Handling And Storage VST (178)	<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 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

**22 Apr 2022 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb VST Audit 07 Handling and stock 2022 attached Please review

**17 Jun 2022 Derek Lamb**

thanks

Audit Goods Inwards and Product Identity  
VST



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 / or dealt with in s	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="#">257393</a> <a href="#">14 Apr 2022</a> 253497	Audit 09 Goods Inward And Product Identity VST (174)	<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 Helen Lamb sent to Derek Lamb

q11 missing info from a srs 68092. think its a system issue

**04 Aug 2022 Derek Lamb**

was a system bug, seem the acount was tagged with a non existant contact ID, not sure why but it wasnt taking the correct number as things didnt match up. fixed.

Non Conformance Issues  
VST

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 / or dealt with in s	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="#">256229</a> <a href="#">05 Apr 2022</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"/>

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

**05 Apr 2022 Helen Lamb**

checked back through all Non Conformance issues to 1st jan 22 nothing of concern or to be investigated or reported on. Order Invoice Error Logs - no issue, nothing to worry about. No feedback issues relating to non conformances that need to me monitored or reviewed

Non Conformance Issues  
VST

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 / or dealt with in s	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="#">250131</a> <a href="#">04 Feb 2022</a>	Non conformance review history VST (286)	<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

**07 Feb 2022 Derek Lamb**

reviewed, nothing stands out as a problem

Audit Picking and Packing  
VST

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 / or dealt with in s						
<a href="#">246776</a> <a href="#">03 Jan 2022</a>	Audit 01 Picking Packing VST (194)	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
<p>Derek Lamb</p> <p>System Generated Audit 01 due 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</p>								
<p><b>04 Mar 2022 Helen Lamb</b></p> <p>Priority Changed From 5 To 2</p>								
<p><b>13 Apr 2022 Helen Lamb</b></p> <p>Next Action Changed From Helen Lamb To Derek Lamb VST Audit completed as per old system as new system not ready Please review the attached.</p>								
<p><b>26 Apr 2022 Derek Lamb</b></p> <p>thankyou</p>								
<p><b>25 Jan 2023 Helen Lamb</b></p> <p>Created Related Issue #284669</p> <p>Added by Helen Lamb sent to Helen Lamb</p> <p>Both Viamed and VSt need a re write as reflects old system</p>								
Non Conformance Issues VST								
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 / or dealt with in s	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="#">244975</a> <a href="#">10 Dec 2021</a> 244667	Audit 17 Internal Audits VST (191) Non Conformance	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
<p>Derek Lamb</p> <p>Added by Derek Lamb sent to Derek Lamb</p> <p>Need a QC21 form to look into this, there should not be blank sections of the standard with regard to an area being audited, unless the section is just the Title of the section. the blanks appearing are define sub sections so no reason to be blank.</p>								
<p><b>13 Dec 2021 Derek Lamb</b></p> <p>sorted</p>								
Non Conformance Issues VST								
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 / or dealt with in s	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="#">241419</a> <a href="#">08 Nov 2021</a>	Shipped Items Return to Supplier BOX828	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
<p>Derek Lamb</p> <p>Does this Return BOX828 warrant a NON conformance report via the CAPA process VM3COP10</p>								
<p><b>10 Nov 2021 Derek Lamb</b></p> <p>VST normal oxygen sensor fails no capa required</p>								
Complaints VST								
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 / or dealt with in s	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="#">239983</a> <a href="#">26 Oct 2021</a>	Objective All complaints to be logged and All complaints to be satisfactorily resolved in 6 months	<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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General

26 Oct 2021 Derek Lamb

No New complaints in the Complaint headers,

Is an Issue/repair with JFD regarding the failure of 1 sensor during a test, not at this time a complaint, but they want to know what the failure is.SRS67874

Audit Management Review

VST

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 / or dealt with in s	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="#">238809</a> <a href="#">15 Oct 2021</a>	Audit 18 Management Review VST (188)	<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 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. 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

**15 Oct 2021 Helen Lamb**

Priority Changed From 5 To 1

**26 Oct 2021 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb audit attached, issues that are outstanding are noted on the viamed audit

**26 Nov 2021 Derek Lamb**

thankyou

Non Conformance Issues

VST

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 / or dealt with in s	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="#">237792</a> <a href="#">05 Oct 2021</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

**07 Oct 2021 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb This is in the process of being built so some functions are not present yet. Most changes to orders are due to the customer making changes. Orders deleted is the highest over the last two months and due to lack of stock- lead times and unpaid proformas. Updates to carriage is the second highest and is due to the customer changing method due to recent delays in shipping by other methods. Covid and Brexit have caused shipping issues recently. No problems stick out and no remedial action is needed on the current figures in the Reasons for opening Orders For Editing page Customer Complaint and Non Conformance Review Screen Nothing since last review re internal errors or areas of concern. Non Conformance Issues - Non Conformance Issues. There is nothing relating to - internal / staff / processes No issues ongoing overall and no problem areas - Viamed

**07 Oct 2021 Derek Lamb**

thank you

Non Conformance Issues

VST

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	Determined Cause of Non Conformity / Complaint	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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		actions required / No Internal Review Requested / Product Failure but no requirement to Escalate Non conformance / or dealt with in s	vigilance Issue requiring a corrective action plan		does not recur	implementation QC 28b	Performance or regulatory requirements	
<a href="#">234773</a> <a href="#">06 Sep 2021</a>	Shipped Items Return to Supplier BOX809	<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 BOX809 warrant a NON conformance report via the CAPA process VM3COP10

**06 Sep 2021 Derek Lamb**

vst Low Output sensor, normal fail tpe no capa required at this point

Audit Analysis of Data  
VST

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 / or dealt with in s	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="#">233619</a> <a href="#">24 Aug 2021</a> 228924	Audit 23 Analysis Of Data VST (185)	<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 Helen Lamb sent to Derek Lamb

Stock Meeting last done april now due

**09 Sep 2021 Derek Lamb**

done

Audit Analysis of Data  
VST

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 / or dealt with in s	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="#">228924</a> <a href="#">13 Jul 2021</a>	Audit 23 Analysis Of Data VST (185)	<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 23 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 Aug 2021 Helen Lamb**

Created Related Issue #233619

Added by Helen Lamb sent to Derek Lamb

Stock Meeting last done april now due

**25 Aug 2021 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb audit done and attached #233619 ongoing issues

**17 Sep 2021 Derek Lamb**

Done

Audit Analysis of Data  
VST

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 / or dealt with in s	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="#">228026</a> <a href="#">05 Jul</a> <a href="#">2021</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

Sales Menu -&gt;

Analysis of Data Viamed Analysis of Data VST

Search for any potential problems in the Graphs provided

Employee Menu Check Audit Roles Titles and Processes in Employee Roles and Titles. 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.

**05 Aug 2021 Derek Lamb**

board meeting held 3 aug 2021 all data was reviewed

Audit Documentation Control

VST

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 / or dealt with in s	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="#">225277</a> <a href="#">10 Jun</a> <a href="#">2021</a>	Audit 10 Documentation Control VST (183)	<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 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

**01 Jul 2021 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb one issue pending related to this 227413. Audit completed and attached

Audit Calibration

VST

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 / or dealt with in s	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="#">223116</a> <a href="#">20 May</a> <a href="#">2021</a>	Audit 06 Calibration VST (182)	<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 06 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

**25 Aug 2021 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb audit done, and attached. No issues outstanding VST doesn't actively use Test equipment as not testing before despatch.

**09 Sep 2021 Derek Lamb**

Done

Audit Analysis of Data

VST

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 / or dealt with in s	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="#">221537</a> <a href="#">05 May 2021</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

Sales Menu -&gt;

Analysis of Data Viamed Analysis of Data VST

Search for any potential problems in the Graphs provided

Employee Menu Check Audit Roles Titles and Processes in Employee Roles and Titles. 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.

**05 May 2021 Derek Lamb**

Helen been on holiday 1 outstanding task overdue, MG on tidying his room, RS Distributor Agreements, t/o dropped in april, need to see how may goes

**Audit Production**

VST

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 / or dealt with in s	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="#">221169</a> <a href="#">03 May 2021</a>	Audit 15 Production VST (175)	<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 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

**04 Aug 2021 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb audit complete some outstanding issue attached to Viamed Audit that relate to both Viamed and VST

**11 Aug 2021 Derek Lamb**

Tasks have been completed

**Audit Post Marketing Surveillance**

VST

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 / or dealt with in s	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="#">221043</a> <a href="#">30 Apr 2021</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"/>
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Derek 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

**01 Jul 2021 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb Audit completed no ongoing issues

**Audit Handling and Storage**

VST

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 / or dealt with in s	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="#">219542</a> <a href="#">15 Apr</a> <a href="#">2021</a>	Audit 07 Handling And Storage VST (178)	<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 07 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 Jun 2021 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb done please see audit attached. No Issues with VST

**15 Jun 2021 Derek Lamb**

thankyou

Non Conformance Issues  
VST

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 / or dealt with in s	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="#">218313</a> <a href="#">06 Apr</a> <a href="#">2021</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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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

**07 Apr 2021 Helen Lamb****13 May 2021 Helen Lamb**

There do not appear to be any inherent problems / frequently occurring issues. When a mistake has been made those responsible have been spoken to and the relevant process re gone over. So the same mistake / error doesn't happen again.

Audit CE Files  
VST

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 / or dealt with in s	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="#">217140</a> <a href="#">22 Mar</a> <a href="#">2021</a>	Audit 12 CE Files VST (176)	<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 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

**01 Jul 2021 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb Audit done no ongoing issues

Audit Goods Inwards and Product Identity  
VST

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 / or dealt with in s	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="#">215995</a> <a href="#">09 Mar</a>	Audit 09 Goods Inward And Product Identity VST	<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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[2021](#) (174)

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

**22 Jun 2021 Helen Lamb**

Created Related Issue #226589

Added by Helen Lamb sent to Derek Lamb

question 16 Quarterly meetings. We do for Viamed, but VST they are annually carried out. Is this ok and correct

**01 Jul 2021 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb one issue pending related to this 220469. Audit completed and attached

#### Non Conformance Issues

VST

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 / or dealt with in s	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="#">212854</a> <a href="#">04 Feb 2021</a>	Non conformance review history VST (286)	<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

**11 Feb 2021 Derek Lamb**

202853, needs closing, when we get to hold a vst meeting

#### Audit Analysis of Data

VST

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 / or dealt with in s	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="#">209745</a> <a href="#">05 Jan 2021</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

Sales Menu ->

Analysis of Data Viamed Analysis of Data VST

Search for any potential problems in the Graphs provided

Employee Menu Check Audit Roles Titles and Processes in Employee Roles and Titles. 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.

**05 Jan 2021 Derek Lamb**

issues system upto date, no old employees, Turnover best on record.

#### Audit Picking and Packing

VST

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 / or dealt with in s	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="#">209457</a> <a href="#">04 Jan</a> <a href="#">2021</a>	Audit 01 Picking Packing VST (194)	<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 01 due 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

**27 Jan 2021 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb audit completed and attached

**27 Jan 2021 Derek Lamb**

thanks no follow on issues

Audit Analysis of Data VST

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 / or dealt with in s	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="#">206996</a> <a href="#">07 Dec</a> <a href="#">2020</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

Sales Menu -&gt;

Analysis of Data Viamed Analysis of Data VST

Search for any potential problems in the Graphs provided

Employee Menu Check Audit Roles Titles and Processes in Employee Roles and Titles. 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 Dec 2020 Derek Lamb**

reviewed, a few issues starting to stack up, but non that require immediate attention, will check again end of week

Audit of Audits VST

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 / or dealt with in s	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="#">206391</a> <a href="#">01 Dec</a> <a href="#">2020</a>	Audit 21 Audit Of Audit VST (192)	<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 21 due Review the Audit Calendar Screen ISO -> Audit Calendar Complete Audit 21 Confirm if Audit calendar needs changing. 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 Oct 2021 Helen Lamb**

audit attached as requested

**13 Oct 2021 Derek Lamb**

thanks

Audit Internal Audits VST

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 / or dealt with in s						
<a href="#">206390</a> <a href="#">01 Dec 2020</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"/>
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. 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								
<b>04 Dec 2020 Helen Lamb</b> Next Action Changed From Helen Lamb To Derek Lamb done								
<b>11 Feb 2021 Derek Lamb</b>								
Audit Analysis of Data VST								
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 / or dealt with in s	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="#">203695</a> <a href="#">05 Nov 2020</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 Sales Menu -> Analysis of Data Viamed Analysis of Data VST Search for any potential problems in the Graphs provided Employee Menu Check Audit Roles Titles and Processes in Employee Roles and Titles. 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.								
<b>05 Nov 2020 Derek Lamb</b> reviewed with bsi last week								
BSI Minor Non conformances VST								
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 / or dealt with in s	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="#">202983</a> <a href="#">29 Oct 2020</a> 155085	vst exclusions	<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 Note VST also now requires an exclusion to 9001 adding will update when bsi approves viameds Next Action Changed From Derek Lamb To Helen Lamb								
<b>29 Oct 2020 Derek Lamb</b> found document in index already 23739 Internal use only VM3COP02.01 Boundaries ISO 9001:2015 VST								
Non Conformance Issues VST								
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 / or dealt with in s	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="#">202853</a> <a href="#">28 Oct</a>	Service Repair Sheet:67494	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>



<b>2020</b>								
Steve Nixon Added by Derek Lamb sent to Derek Lamb <b>Service Repair Sheet:67494</b> SRN32620 Oxygen Sensor VST 8010050 100587 Oxygen Sensor - JFD (non-mag).. Account Non Conformance should have been raised against the sensors with incorrect body size								
<b>28 Oct 2020 Derek Lamb</b>								
Header Changed From 202 VIAMED Stock Repairs Review - General To VST Management Non Conformance Issues								
<b>28 Oct 2020 Derek Lamb</b>								
Next Action Changed From Derek Lamb To Steve Nixon Noted its Envitec Issue, however its as close to a customer complaint, via a product return. as the size was incorrect, Due to being in covid lock down and the urgency to get these turned around, a non conformance was not raised. As we do not test the sensors from Envitec we will not catch these in the future, could do with finding a response from Envitec as to what they have done to stop this happening again. We may already have this in the system, but it need printing and attaching to this issue.								
<b>15 Nov 2021 Derek Lamb</b>								
see srs 67494								
Audit Management Review								
VST								
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 / or dealt with in s	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="#">20080115 Oct 2020</a>	Audit 18 Management Review VST (188)	<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 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. 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								
<b>19 Oct 2020 Helen Lamb</b>								
Next Action Changed From Helen Lamb To Derek Lamb done please review this and check you are happy with it								
<b>11 Feb 2021 Derek Lamb</b>								
thanks								
Non Conformance Issues								
VST								
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 / or dealt with in s	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="#">19946105 Oct 2020</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"/>
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								
<b>09 Oct 2020 Helen Lamb</b>								
No errors reoccurring								
Audit Analysis of Data								
VST								
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 / or dealt with in s	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="#">199451</a> <a href="#">05 Oct</a> <a href="#">2020</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"/>
<p>Derek Lamb System Generated</p> <p>Sales Menu -&gt;</p> <p>Analysis of Data Viamed Analysis of Data VST</p> <p>Search for any potential problems in the Graphs provided</p> <p>Employee Menu Check Audit Roles Titles and Processes in Employee Roles and Titles. 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.</p> <p><b>07 Oct 2020 Derek Lamb</b> Check Audit Roles Titles and Processes in Employee Roles and Titles. Done, updated to make easier mass closing of relevant issues if there is a back log , and the backlog reason is fixed (i.e. covid build up non essential jobs) While resources have been tight personnel wise, we have show we can deal with a sudden large through put of stock Issue statistics now broken,</p>								
Non Conformance Issues VST								
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 / or dealt with in s	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="#">198689</a> <a href="#">25 Sep</a> <a href="#">2020</a>	Shipped Items Return to Supplier BOX741	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
<p>Derek Lamb Does this Return BOX741 warrant a NON conformance report via the CAPA process VM3COP10</p> <p><b>28 Sep 2020 Derek Lamb</b> VST sensors, no cap required</p>								
Non Conformance Issues VST								
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 / or dealt with in s	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="#">172324</a> <a href="#">06 Apr</a> <a href="#">2020</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"/>
<p>Helen Lamb System Generated Check the history of the last Non conformance review,</p> <p>check actions are being carried out, and non conformances are not reoccurring</p> <p><b>23 Apr 2020 Helen Lamb</b> nothing on going for VST</p>								
Any New QC21 Forms VST								
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 / or dealt with in s	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="#">167500</a> <a href="#">03 Mar</a> <a href="#">2020</a>	QC 21 Form Oxygen Sensor - R17JJ-CCR Shipment Dates	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
<p>Derek Lamb Added by Derek Lamb sent to Derek Lamb JJCCR usually place a forward order, and need reminding when to place another order, this month the reminder was late, and while we have stock in the pipeline, Jan will usually want the sensors at the start of the month, Our required date on next supplier order was set as the 4th March, When the confirmation came in on the</p>								

3rd feb, and the estimate shipping date correctly filled in it was for the date 11th March. the dates mismatch was not picked up by office staff while confirming the order

**03 Mar 2020 Derek Lamb**

Raised a QC21 Form due to the fact the customer is un happy, and his stock is going to be a week later than planned

**11 Mar 2020 Derek Lamb**

Created Related Issue #168340

Added by Derek Lamb sent to Catrin Hird

**09 Apr 2020 Derek Lamb**

Header Changed From 503 VST Management Non Conformance Issues To Non Conformance Issues Any New QC21 Forms

Urgent Flag Changed To On

Subject Changed From Oxygen Sensor - R17JJ-CCR Shipment Dates

Subject Changed To QC 21 Form Oxygen Sensor - R17JJ-CCR Shipment Dates

**10 Apr 2020 Derek Lamb**

### Non Conformance Issues VST

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 / or dealt with in s	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="#">164203</a> <a href="#">04 Feb 2020</a>	Non conformance review history VST (286)	<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

**12 Mar 2020 Derek Lamb**

upto date as feb

### Audit Picking and Packing VST

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 / or dealt with in s	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="#">161466</a> <a href="#">02 Jan 2020</a>	Audit 01 Picking Packing VST (194)	<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 01 due 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

**08 Jan 2020 Helen Lamb**

Priority Changed From 5 To 2

**10 Sep 2020 Helen Lamb**

Created Related Issue #196533

Added by Helen Lamb sent to Sarah Walton

Order ID RST124142 has no customer paperwork attached. Please can you find the paperwork and add it also please make sure to add customer paperwork to all orders.

**23 Sep 2020 Helen Lamb**

Next Action Changed From Helen Lamb To Derek Lamb Audit completed and attached, please check you are happy with this and let me know if you have any queries.

**28 Sep 2020 Derek Lamb**

checked and ok

Audit Design Control

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 / or dealt with in s	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="#">161465</a> <a href="#">02 Jan 2020</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"/>
<p>Derek Lamb</p> <p>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</p>								
<p><b>08 Jan 2020 Helen Lamb</b></p> <p>Priority Changed From 5 To 2</p>								
<p><b>23 Sep 2020 Helen Lamb</b></p> <p>Next Action Changed From Helen Lamb To Derek Lamb Audit completed and attached, please check you are happy with this and let me know if you have any queries.</p>								
<p><b>28 Sep 2020 Derek Lamb</b></p> <p>checked ok, no viamed design,</p>								