

Quality Management System Route Map to Documents and Procedures

VST Ltd ISO9001:2015

Version Date: 13 Dec 2021

Listing of Current Sections

Section	Documents related	Processes Direct Links
4 Context of the organization		
4 Context of the organization	Top Level Document: QMS Route Map VST Ltd ISO9001_2015 Revision Document ID75598 **Date Revision 22 Nov 2021 Reviewed 22 Nov 2021 Top Level Document: Need Risks and Expectations of External Parties VST Revision Document ID74925 Date Revision 15 Nov 2021 Reviewed 15 Nov 2021 Top Level Document: VST ISO 9001:2015 Scope Revision Document ID24442 Date Revision 01 Dec 2017 Reviewed 27 Sep 2021 Chart 39 external parties vst Revision Document ID22630 Date Revision 14 Oct 2017 Reviewed 14 Oct 2017 BS EN ISO 9001:2015 Revision Document ID16229 Date Revision 01 Feb 2016 Reviewed 01 Feb 2016	Process: 7433 Responsibility Allocation : VST Board Directors Meeting 09 Mar 2016

	Chart 43 Processes and Intrastats Revision Document ID23561 Date Revision 28 Oct 2017 Reviewed 28 Oct 2017 Chart 42 Processes, Tasks and Audits Review Revision Document ID23559 Date Revision 28 Oct 2017 Reviewed 28 Oct 2017 Chart 40 Management review plan Issues followup Revision Document ID22458 Date Revision 05 Oct 2017 Reviewed 05 Oct 2017 VM3COP24.01 Definitions of Risk Revision Document ID75525 Date Revision 19 Nov 2021 Reviewed 19 Nov 2021 Intrastats overview Revision Document ID23567 Date Revision 28 Oct 2017 Reviewed 28 Oct 2017 VST TOP Level Objectives Revision Document ID46732 Date Revision 29 Oct 2020 Reviewed 29 Oct 2020 VM3COP02.01 Boundaries ISO 9001:2015 VST Revision Document ID69692 Date Revision 14 Sep 2021 Reviewed 14 Sep 2021	
4.1 The organization shall determine external and internal issues that are relevant to its purpose and its strategic direction and that affect its ability to achieve the intended result(s) of its quality management system.	Top Level Document: VOP 24 Needs, Risks and Expectations of External Parties Revision Document ID31060 Date Revision 30 Sep 2019 Reviewed 30 Sep 2019 Top Level Document: Need Risks and Expectations of External Parties VST	Process: 7837 Review External Parties Influencing The QMS VST / Viamed 23 Sep 2017

The organization shall monitor and review information about these external and internal issues. NOTE 1 Issues can include positive and negative factors or conditions for consideration. NOTE 2 Understanding the external context can be facilitated by considering issues arising from legal, technological, competitive, market, cultural, social and economic environments, whether international, national, regional or local. NOTE 3 Understanding the internal context can be facilitated by considering issues related to values, culture, knowledge and performance of the organization. Understanding the organization and its context	Revision Document ID74925 Date Revision 15 Nov 2021 Reviewed 15 Nov 2021 Audit 18 Management Review Revision Document ID73320 **Date Revision 26 Oct 2021 Reviewed 26 Oct 2021 Chart 39 external parties vst Revision Document ID22630 Date Revision 14 Oct 2017 Reviewed 14 Oct 2017	
4.2 Due to their effect or potential effect on the organization's ability to consistently provide products and services that meet customer and applicable statutory and regulatory requirements, the organization shall determine: a) the interested parties that are relevant to the quality management system; b) the requirements of these interested parties that are relevant to the quality management system. The organization shall monitor and review information about these interested parties and their relevant requirements. Understanding the needs and expectations of interested parties	Top Level Document: Need Risks and Expectations of External Parties VST Revision Document ID74925 Date Revision 15 Nov 2021 Reviewed 15 Nov 2021 Top Level Document: VOP 24 Needs, Risks and Expectations of External Parties Revision Document ID31060 Date Revision 30 Sep 2019 Reviewed 30 Sep 2019 Audit 18 Management Review Revision Document ID73320 **Date Revision 26 Oct 2021 Reviewed 26 Oct 2021 Chart 39 external parties vst Revision Document ID22630 Date Revision 14 Oct 2017 Reviewed 14 Oct 2017	Process: 7792 Shipped Order Success Report 13 Mar 2017 Process: 7740 Weights Per Region Needed To Submit EC Sales List 13 Sep 2016 Process: 7734 Responsibility Allocation : Humanmed Order Processing 25 Aug 2016 Process: 7710 Responsibility Allocation : Proforma And Quote Processing 29 Jun 2016 Process: 7709 Delivered not Invoiced 28 Jun 2016 Process: 7953 Vandagraph Delivery Notifications 26 May 2020 Process: 7691 Ship Sale Or Returns 21 Apr 2016 Process: 7690 Ship Repairs 21 Apr 2016 Process: 7686

Thorough Checking Of Awaiting Action Tray 21 Apr 2016

Process: 7685

Repairs Ready For Invoice 18 Apr 2016

Process: 7684

Repairs Ready For Quote 18 Apr 2016

Process: 7678

Check Catalog 360 Circle For Quotes And Orders 08 Apr 2016

Process: 7674

Check Repairs Ready For Invoice List 10 Mar 2016

Process: 7673

Check Expiry Dated Stock 09 Mar 2016

Process: 7398

Responsibility Allocation : VST Stock Meeting UPS Shipping Fuel
Surcharge 09 Mar 2016

Process: 7396

Responsibility Allocation : VST Stock Meeting `Goods Out` Review
09 Mar 2016

Process: 7394

Responsibility Allocation : VST Stock Meeting Repairs Review -
General 09 Mar 2016

Process: 7388

Responsibility Allocation : VST Stock Meeting Returns Overview 09
Mar 2016

Process: 7837

Review External Parties Influencing The QMS VST / Viamed 23 Sep
2017

Process: 7385

Responsibility Allocation : VST Stock Meeting Sales Forward Orders
Review 09 Mar 2016

Process: 6956

Responsibility Allocation : Sales Order Issues 09 Mar 2016

Process: 7090

Responsibility Allocation : Office Procedures 09 Mar 2016

Process: 6938

Responsibility Allocation : Customer Database Updates 09 Mar 2016

Process: 2

Answering Telephones 16 Feb 2016

Process: 5

Responsibility Allocation : Processing Of Sales Orders 16 Feb 2016

Process: 6

Responsibility Allocation : Updating Contact Management System 16 Feb 2016

Process: 7

Responsibility Allocation : Checking Of Sales Orders 16 Feb 2016

Process: 8

Responsibility Allocation : Order And Status Liaison With Customers 16 Feb 2016

Process: 6898

GHX Web Pricing 09 Mar 2016

Process: 5876

E.Commerce Cardea And Multiquote 17 Feb 2016

Process: 5871

Check Sale Or Returns 17 Feb 2016

Process: 9

Distribution Of Faxes 16 Feb 2016

Process: 10

Distribution Of Emails 16 Feb 2016

Process: 11

Distribution Of Mail 16 Feb 2016

Process: 14

Fax Paper 16 Feb 2016

Process: 15

Filing and Archiving 16 Feb 2016

Process: 16

Responsibility Allocation : Photocopying 16 Feb 2016

Process: 21

Office Sales Projects 16 Feb 2016

Process: 36

Emailing Of Invoices 16 Feb 2016

Process: 5879

Responsibility Allocation : Customer Returning Goods On Our UPS Account 18 Feb 2016

Process: 5882

Responsibility Allocation : Send Post To Humanmed 24 Feb 2016

Process: 5891

Processing Of Repair Quotes And Orders 25 Feb 2016

Process: 5892

Checking EBay And Amazon For Orders And Messages 25 Feb 2016

Process: 5893

Answering Website Questions 25 Feb 2016

Process: 5894

Checking Of Active List 25 Feb 2016

Process: 5895

Responsibility Allocation : Completing Office Job List 25 Feb 2016

Process: 5896

Responsibility Allocation : Ensuring ORD's Are Taken To Goods Out And Invoices Are Retrieved 25 Feb 2016

Process: 5899

Proforma And Quote Chasing 25 Feb 2016

Process: 5901

Link Call Log Contacts To The CRM 02 Mar 2016

Process: 5913

Check For Humanmed Orders In Logistics Mailbox 03 Mar 2016

Process: 5943

Check Cardea And Multiquote 08 Mar 2016

Process: 5944

Responsibility Allocation : Chasing Lost Customers 08 Mar 2016

Process: 5945

Responsibility Allocation : Sending Samples 08 Mar 2016

Process: 5946

Responsibility Allocation : Sending Sale Or Returns 08 Mar 2016

Process: 7693

Collect Repair Filing From Warehouse 22 Apr 2016

Process: 5948

Adding New Accounts To Opera 08 Mar 2016

Process: 5949

Filling Credit Card Slips 08 Mar 2016

Process: 6958

Responsibility Allocation : Shipped Order Queries 09 Mar 2016

Process: 7676

PDFing Of Invoices Viamed 17 Mar 2016

Process: 7699

Shred Sensitive Paperwork In JL Office 19 May 2016

Process: 7712

Review Inward Payments 01 Jul 2016

Process: 7735

Ensure SOR's Are Followed Up 01 Sep 2016

		Process: 7752 SRS Folder 22 Nov 2016 Process: 7758 Check For GHX Orders 17 Jan 2017 Process: 7760 Send Service Offers 31 Jan 2017 Process: 7761 Send VST Delivery Notifications 01 Feb 2017 Process: 7783 PDF VST Invoices And Purchase Orders 10 Feb 2017 Process: 7795 Answering UK Web Questions 27 Apr 2017 Process: 6954 Back Orders Review - By Customer 09 Mar 2016 Process: 5859 Review Un-shipped Parcels 17 Feb 2016 Process: 7748 Check Repair Orders 10 Oct 2016 Process: 7749 Check Repair Quotes 10 Oct 2016 Process: 5872 Check Sale Or Returns Export 17 Feb 2016 Process: 5875 Check Paypal For Orders 17 Feb 2016
4.3 The organization shall determine the boundaries and applicability of the quality management system to establish its scope. When determining this scope, the organization shall consider: a) the external and internal issues referred to in 4.1; b) the requirements of relevant interested parties referred to in 4.2; c) the products and services of the organization. The organization shall apply all the requirements of this International Standard	Top Level Document: VOP 01 Documentation and Records, Control, Creation, Storage, Retrieval, Revision Control and Online Records Revision Document ID75407 Date Revision 18 Nov 2021 Reviewed 18 Nov 2021 Top Level Document: VST ISO 9001:2015 Scope Revision Document ID24442 Date Revision 01 Dec 2017 Reviewed 27 Sep 2021 Audit 18 Management Review Revision Document ID73320 **Date Revision 26 Oct 2021 Reviewed	Process: 7744 FDA Device Establishment Registration And Listing 28 Sep 2016 Process: 7668 Responsibility Allocation : Upgrading Intrastats ISO Quality system 09 Mar 2016 Process: 7389 Responsibility Allocation : VST Stock Meeting Returns Overview - From Customers 09 Mar 2016 Process: 7837 Review External Parties Influencing The QMS VST / Viamed 23 Sep 2017 Process: 7848 Review ISO Scopes 27 Sep 2017 Process: 7871

if they are applicable within the determined scope of its quality management system. The scope of the organization's quality management system shall be available and be maintained as documented information. The scope shall state the types of products and services covered, and provide justification for any requirement of this International Standard that the organization determines is not applicable to the scope of its quality management system. Conformity to this International Standard may only be claimed if the requirements determined as not being applicable do not affect the organization's ability or responsibility to ensure the conformity of its products and services and the enhancement of customer satisfaction. Determining the scope of the quality management system	26 Oct 2021 VM3COP02.01 Boundaries ISO 9001:2015 VST Revision Document ID69692 Date Revision 14 Sep 2021 Reviewed 14 Sep 2021	Review Exclusion From Viamed 13485:2016 And VST 9001:2015 15 Oct 2017
4.4 Quality management system and its processes	Top Level Document: QMS Route Map VST Ltd ISO9001_2015 Revision Document ID75598 **Date Revision 22 Nov 2021 Reviewed 22 Nov 2021	
4.4.1 The organization shall establish, implement, maintain and continually improve a quality management system, including the processes needed and their interactions, in accordance with the requirements of this International Standard. The organization shall determine the	Audit 10 Documentation Control Revision Document ID63807 Date Revision 30 Jun 2021 Reviewed 30 Jun 2021 Audit 20 Process verification to Managment Revision Document ID73324 Date Revision 26 Oct 2021 Reviewed 26 Oct 2021 Chart 34 Process Teams Org Chart	Process: 7837 Review External Parties Influencing The QMS VST / Viamed 23 Sep 2017

processes needed for the quality management system and their application throughout the organization, and shall:

- a) determine the inputs required and the outputs expected from these processes;
- b) determine the sequence and interaction of these processes;
- c) determine and apply the criteria and methods (including monitoring, measurements and related performance indicators) needed to ensure the effective operation and control of these processes;
- d) determine the resources needed for these processes and ensure their availability;
- e) assign the responsibilities and authorities for these processes;
- f) address the risks and opportunities as determined in accordance with the requirements of 6.1;
- g) evaluate these processes and implement any changes needed to ensure that these processes achieve their intended results;
- h) improve the processes and the quality management system

Revision Document ID8707

Date Revision 12 Oct 2011 Reviewed 12 Oct 2011

Chart 33 Launch of a new product

Revision Document ID8706

Date Revision 12 Oct 2011 Reviewed 12 Oct 2011

Employee Roles

Revision Document ID20125

Date Revision 16 May 2017 Reviewed 16 May 2017

Employee roles Example Process

Revision Document ID20129

Date Revision 16 May 2017 Reviewed 16 May 2017

Employee Roles Individual Processes

Revision Document ID20127

Date Revision 16 May 2017 Reviewed 16 May 2017

Explanation Employee Roles and Titles

Revision Document ID22144

Date Revision 20 Sep 2017 Reviewed 20 Sep 2017

Explanation Employee Roles Titles

Responsibilities Processes and Repeating Tasks Monitoring

Revision Document ID22287

Date Revision 27 Sep 2017 Reviewed 27 Sep 2017

Chart 32 Generic Sales Process

Revision Document ID8705

Date Revision 12 Oct 2011 Reviewed 12 Oct 2011

Chart 31 Chart Interfaces

Revision Document ID8704

Date Revision 12 Oct 2011 Reviewed 12 Oct 2011

Chart 30 System Design Plan

Revision Document ID8703

Date Revision 12 Oct 2011 Reviewed 12 Oct 2011

Chart 29 Sales Acquisition

Revision Document ID8702

Date Revision 12 Oct 2011 Reviewed 12 Oct 2011

Chart 28 Quarantine and Hold

Revision Document ID8701

Date Revision 12 Oct 2011 Reviewed 12 Oct 2011

Chart 27 Customer Complaints Chart 27

Revision Document ID8700

Date Revision 12 Oct 2011 Reviewed 12 Oct 2011

Chart 26 Data Analysis

Revision Document ID8699

Date Revision 12 Oct 2011 Reviewed 12 Oct 2011

Chart 25 Inspection and Test

Revision Document ID8698

Date Revision 12 Oct 2011 Reviewed 12 Oct 2011

Chart 24 Goods Inwards

Revision Document ID8697

Date Revision 12 Oct 2011 Reviewed 12 Oct 2011

Chart 23 Picking and Packing

Revision Document ID8696

Date Revision 12 Oct 2011 Reviewed 12 Oct 2011

Chart 22 Stock Control

Revision Document ID8695

Date Revision 12 Oct 2011 Reviewed 12 Oct 2011

Chart 21 Repairs

Revision Document ID8694

Date Revision 12 Oct 2011 Reviewed 12 Oct 2011

Chart 20 Production

Revision Document ID8693

Date Revision 12 Oct 2011 Reviewed 12 Oct 2011

Chart 19 HSE Risk Assesments

Revision Document ID8692

Date Revision 12 Oct 2011 Reviewed 12 Oct 2011

Chart 18 Calibration

Revision Document ID8691

Date Revision 12 Oct 2011 Reviewed 12 Oct 2011

Chart 17 Design Repairs

Revision Document ID8690

Date Revision 12 Oct 2011 Reviewed 12 Oct 2011

Chart 16 Internal Audits

Revision Document ID8689

Date Revision 12 Oct 2011 Reviewed 12 Oct 2011

Chart 15 Purchasing

Revision Document ID8688

Date Revision 12 Oct 2011 Reviewed 12 Oct 2011

Chart 13 Sales Orders

Revision Document ID8687

Date Revision 12 Oct 2011 Reviewed 12 Oct 2011

Chart 12 Infrastructure and Environment

Revision Document ID8686

Date Revision 12 Oct 2011 Reviewed 12 Oct 2011

Chart 11 Provision of Resources

Revision Document ID8685

Date Revision 12 Oct 2011 Reviewed 12 Oct 2011

Chart 10 Documentation

Revision Document ID8684

Date Revision 12 Oct 2011 Reviewed 12 Oct 2011
Chart 09 Management System
Revision Document ID8683
Date Revision 12 Oct 2011 Reviewed 12 Oct 2011
Chart 08 Correction and Prevention
Revision Document ID8682
Date Revision 12 Oct 2011 Reviewed 12 Oct 2011
Chart 07 Measurement and Analysis
Revision Document ID8681
Date Revision 12 Oct 2011 Reviewed 12 Oct 2011
Chart 06 General Process Control
Revision Document ID8680
Date Revision 12 Oct 2011 Reviewed 12 Oct 2011
Chart 05 Product Realisation
Revision Document ID8679
Date Revision 12 Oct 2011 Reviewed 12 Oct 2011
Chart 04 Design and Development
Revision Document ID8678
Date Revision 12 Oct 2011 Reviewed 12 Oct 2011
Chart 03 Customer Requirements
Revision Document ID8677
Date Revision 12 Oct 2011 Reviewed 12 Oct 2011
Chart 02 Resource Management
Revision Document ID8676
Date Revision 12 Oct 2011 Reviewed 12 Oct 2011
Chart 01 System and Documentation
Revision Document ID8675
Date Revision 12 Oct 2011 Reviewed 12 Oct 2011
Chart 00 System Model

	Revision Document ID8674 Date Revision 12 Oct 2011 Reviewed 12 Oct 2011	
4.4.2 To the extent necessary, the organization shall: a) maintain documented information to support the operation of its processes; b) retain documented information to have confidence that the processes are being carried out as planned.	Top Level Document: VOP 01 Documentation and Records, Control, Creation, Storage, Retrieval, Revision Control and Online Records Revision Document ID75407 Date Revision 18 Nov 2021 Reviewed 18 Nov 2021 Audit 10 Documentation Control Revision Document ID63807 Date Revision 30 Jun 2021 Reviewed 30 Jun 2021 4.4.2 Quality management system and its processes Revision Document ID22132 Date Revision 20 Sep 2017 Reviewed 20 Sep 2017	Process: 7713 Review Roles And Responsibilitys 17 Aug 2016 Process: 27 Management Reviews And Quality Audits 16 Feb 2016 Process: 7705 Checking For Uploaded Files 08 Jun 2016 Process: 7693 Collect Repair Filing From Warehouse 22 Apr 2016 Process: 7692 Responsibility Allocation : Take Complete Repair Paperwork To Office 22 Apr 2016

5 Leadership

5 Leadership		
5.1 Leadership and commitment		
5.1.1 Top management shall demonstrate leadership and commitment with respect to the quality management system by: a) taking accountability for the effectiveness of the quality management system; b) ensuring that the quality policy and quality objectives are established for the quality management	Top Level Document: VM3COP00.00 Viamed Quality Statement policy and objectives Revision Document ID22684 Date Revision 16 Oct 2017 Reviewed 03 Aug 2021 Top Level Document: VM3COP02.02 Viamed Company Responsibilitys organisation chart structure Revision Document ID27474 Date Revision 20 Sep 2018 Reviewed 03	Process: 22 Company Polycys 16 Feb 2016 Process: 23 Company Objectives 16 Feb 2016 Process: 26 Company Resources 16 Feb 2016 Process: 7834 Financial Review 20 Sep 2017 Process: 27 Management Reviews And Quality Audits 16 Feb 2016 Process: 7750

system and are compatible with the context and strategic direction of the organization;

c) ensuring the integration of the quality management system requirements into the organization's business processes;

d) promoting the use of the process approach and risk-based thinking;

e) ensuring that the resources needed for the quality management system are available;

f) communicating the importance of effective quality management and of conforming to the quality management system requirements;

g) ensuring that the quality management system achieves its intended results;

h) engaging, directing and supporting persons to contribute to the effectiveness of the quality management system;

i) promoting improvement;

j) supporting other relevant management roles to demonstrate their leadership as it applies to their areas of responsibility.

NOTE Reference to "business" in this International Standard can be interpreted broadly to mean those activities that are core to the purposes of the organization's existence, whether the organization is public, private, for profit or not for profit. **General**

Aug 2021

Top Level Document: VOP 02 Personnel and Responsibility , Staff and Staffing Issues, Training, Roles and Tasks

Revision Document ID73529

Date Revision 29 Oct 2021 Reviewed 29 Oct 2021

Top Level Document: VOP 18 Maintenance Building, Fabric and Infrastructure

Revision Document ID31036

Date Revision 30 Sep 2019 Reviewed 30 Sep 2019

Audit 08 Training, Competence and Human Resources

Revision Document ID70147

Date Revision 20 Sep 2021 Reviewed 20 Sep 2021

Audit 10 Documentation Control

Revision Document ID63807

Date Revision 30 Jun 2021 Reviewed 30 Jun 2021

VM3COP00.00 VST Quality Statement policy and objectives

Revision Document ID22062

Date Revision 16 Sep 2017 Reviewed 24 Aug 2021

VM3COP02 Organisation

Responsibilities Viamed

Revision Document ID17423

Date Revision 07 Sep 2016 Reviewed 07 Sep 2016

Explanation Quality Objectives

Revision Document ID18483

Date Revision 18 Jan 2017 Reviewed 18 Jan 2017

Audit 18 Management Review

Revision Document ID73320

****Date Revision 26 Oct 2021 Reviewed**

Meeting With Management 14 Oct 2016

Process: 7753

Management Meeting Warehouse 22 Nov 2016

Process: 7093

BSI Audits Calander 09 Mar 2016

Process: 7739

Intrastats Amendment Log 12 Sep 2016

Process: 7743

Customer Complaints Paper File 26 Sep 2016

Process: 6931

Customer Complaints 09 Mar 2016

Process: 7833

Importance Of Effective Quality Management 20 Sep 2017

Process: 7199

Non Conformities Review Viamed 09 Mar 2016

Process: 7828

Review The Quality Policy Viamed 16 Sep 2017

Process: 7827

Review The Quality Policy VST 16 Sep 2017

Process: 7791

Price List Check 10 Mar 2017

Process: 7744

FDA Device Establishment Registration And Listing 28 Sep 2016

Process: 7697

Yearly Pricing Review 09 May 2016

Process: 7670

Humanmed general Issues 09 Mar 2016

Process: 7668

Responsibility Allocation : Upgrading Intrastats ISO Quality system 09 Mar 2016

26 Oct 2021

**Audit 20 Process verification to
Managment**

Revision Document ID73324

Date Revision 26 Oct 2021 Reviewed 26
Oct 2021

Explanation Control of documents

Revision Document ID21322

Date Revision 06 Aug 2017 Reviewed 06
Aug 2017

VM3COP19 Health and Safety

Revision Document ID21800

Date Revision 05 Sep 2017 Reviewed 05
Sep 2017

Explanation Employee Roles and Titles

Revision Document ID22144

Date Revision 20 Sep 2017 Reviewed 20
Sep 2017

Viamed Top Level Quality Objectives

Revision Document ID22429

Date Revision 04 Oct 2017 Reviewed 04
Oct 2017

**Chart 40 Management review plan
Issues followup**

Revision Document ID22458

Date Revision 05 Oct 2017 Reviewed 05
Oct 2017

Chart 01 System and Documentation

Revision Document ID8675

Date Revision 12 Oct 2011 Reviewed 12
Oct 2011

Chart 02 Resource Management

Revision Document ID8676

Date Revision 12 Oct 2011 Reviewed 12
Oct 2011

How to Hold Intrastat Meetings

Revision Document ID8928

Date Revision 18 Oct 2011 Reviewed 18
Oct 2011

	VM3COP24.01 Definitions of Risk Revision Document ID75525 Date Revision 19 Nov 2021 Reviewed 19 Nov 2021	
5.1.2 5.1.2 Customer focus Top management shall demonstrate leadership and commitment with respect to customer focus by ensuring that: a) customer and applicable statutory and regulatory requirements are determined, understood and consistently met; b) the risks and opportunities that can affect conformity of products and services and the ability to enhance customer satisfaction are determined and addressed; c) the focus on enhancing customer satisfaction is maintained. Customer focus	Top Level Document: VOP 07 Stock Control, Handling, Control of Labelling, Storage, Movement Revision Document ID75973 **Date Revision 24 Nov 2021 Reviewed 24 Nov 2021 Top Level Document: VOP 19 Feedback Customer Complaints Vigilance and Notifications Viamed Ltd Revision Document ID75475 Date Revision 18 Nov 2021 Reviewed 18 Nov 2021 Top Level Document: VOP 03 Contract Review, Enquires, Office Processes Revision Document ID75949 **Date Revision 24 Nov 2021 Reviewed 24 Nov 2021 Audit 16 Sales and Marketing Revision Document ID69457 Date Revision 10 Sep 2021 Reviewed 10 Sep 2021 Audit 02 Contract Review and Sales Order Processing Revision Document ID69328 Date Revision 09 Sep 2021 Reviewed 09 Sep 2021 Audit 01 Picking packing Revision Document ID51629 Date Revision 13 Jan 2021 Reviewed 13 Jan 2021 Audit 02 Contract Review and Sales Order Processing Revision Document ID69328 Date Revision 09 Sep 2021 Reviewed 09 Sep 2021	Process: 7830 Review Q.A. Failures Report 18 Sep 2017 Process: 7825 Responsibility Allocation : Order Picking 06 Sep 2017 Process: 7822 Review Oxylink Stock 26 Jul 2017 Process: 7801 VST Price Review 17 May 2017 Process: 7797 Check Order Are Being Picked In Priority Order 10 May 2017 Process: 7791 Price List Check 10 Mar 2017 Process: 7761 Send VST Delivery Notifications 01 Feb 2017 Process: 7758 Check For GHX Orders 17 Jan 2017 Process: 7735 Ensure SOR's Are Followed Up 01 Sep 2016 Process: 7734 Responsibility Allocation : Humanmed Order Processing 25 Aug 2016 Process: 7710 Responsibility Allocation : Proforma And Quote Processing 29 Jun 2016 Process: 7709 Delivered not Invoiced 28 Jun 2016 Process: 7697 Yearly Pricing Review 09 May 2016 Process: 7953 Vandagraph Delivery Notifications 26 May 2020 Process: 7691 Ship Sale Or Returns 21 Apr 2016 Process: 7690 Ship Repairs 21 Apr 2016 Process: 7686

VM3COP20.01 Post In Distributing the Post

Revision Document ID18641

Date Revision 10 Feb 2017 Reviewed 10 Feb 2017

VM3COP10.02 Product Recall locate products out in the Field

Revision Document ID74788

Date Revision 12 Nov 2021 Reviewed 12 Nov 2021

Thorough Checking Of Awaiting Action Tray 21 Apr 2016

Process: 7685

Repairs Ready For Invoice 18 Apr 2016

Process: 7684

Repairs Ready For Quote 18 Apr 2016

Process: 7683

Check Stock For Proforma 18 Apr 2016

Process: 7678

Check Catalog 360 Circle For Quotes And Orders 08 Apr 2016

Process: 7676

PDFing Of Invoices Viamed 17 Mar 2016

Process: 7674

Check Repairs Ready For Invoice List 10 Mar 2016

Process: 7673

Check Expiry Dated Stock 09 Mar 2016

Process: 7670

Humanmed general Issues 09 Mar 2016

Process: 7398

Responsibility Allocation : VST Stock Meeting UPS Shipping Fuel Surcharge 09 Mar 2016

Process: 7396

Responsibility Allocation : VST Stock Meeting `Goods Out` Review 09 Mar 2016

Process: 7394

Responsibility Allocation : VST Stock Meeting Repairs Review - General 09 Mar 2016

Process: 7390

Responsibility Allocation : VST Stock Meeting Returns Overview - Credits 09 Mar 2016

Process: 7389

Responsibility Allocation : VST Stock Meeting Returns Overview - From Customers 09 Mar 2016

Process: 7385

Responsibility Allocation : VST Stock Meeting Sales Forward Orders Review 09 Mar 2016

Process: 6938

Responsibility Allocation : Customer Database Updates 09 Mar 2016

Process: 6956

Responsibility Allocation : Sales Order Issues 09 Mar 2016

Process: 5871

Check Sale Or Returns 17 Feb 2016

Process: 5876

E.Commerce Cardea And Multiquote 17 Feb 2016

Process: 6898

GHX Web Pricing 09 Mar 2016

Process: 7090

Responsibility Allocation : Office Procedures 09 Mar 2016

Process: 5872

Check Sale Or Returns Export 17 Feb 2016

Process: 2

Answering Telephones 16 Feb 2016

Process: 5

Responsibility Allocation : Processing Of Sales Orders 16 Feb 2016

Process: 6

Responsibility Allocation : Updating Contact Management System 16 Feb 2016

Process: 7

Responsibility Allocation : Checking Of Sales Orders 16 Feb 2016

Process: 8

Responsibility Allocation : Order And Status Liaison With Customers 16 Feb 2016

Process: 9

Distribution Of Faxes 16 Feb 2016

Process: 10

Distribution Of Emails 16 Feb 2016

Process: 11

Distribution Of Mail 16 Feb 2016

Process: 14

Fax Paper 16 Feb 2016

Process: 15

Filing and Archiving 16 Feb 2016

Process: 16

Responsibility Allocation : Photocopying 16 Feb 2016

Process: 21

Office Sales Projects 16 Feb 2016

Process: 36

Emailing Of Invoices 16 Feb 2016

Process: 5879

Responsibility Allocation : Customer Returning Goods On Our UPS Account 18 Feb 2016
Process: 5875
Check Paypal For Orders 17 Feb 2016
Process: 5882
Responsibility Allocation : Send Post To Humanmed 24 Feb 2016
Process: 5891
Processing Of Repair Quotes And Orders 25 Feb 2016
Process: 5892
Checking EBay And Amazon For Orders And Messages 25 Feb 2016
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Process: 5895
Responsibility Allocation : Completing Office Job List 25 Feb 2016
Process: 5896
Responsibility Allocation : Ensuring ORD's Are Taken To Goods Out And Invoices Are Retrieved 25 Feb 2016
Process: 5899
Proforma And Quote Chasing 25 Feb 2016
Process: 5901
Link Call Log Contacts To The CRM 02 Mar 2016
Process: 5913
Check For Humanmed Orders In Logistics Mailbox 03 Mar 2016
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Responsibility Allocation : Chasing Lost Customers 08 Mar 2016
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Process: 5946
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Process: 5948
Adding New Accounts To Opera 08 Mar 2016
Process: 5949
Filling Credit Card Slips 08 Mar 2016
Process: 5947
Responsibility Allocation : Search For Distributors 08 Mar 2016

Process: 6958

Responsibility Allocation : Shipped Order Queries 09 Mar 2016

Process: 7693

Collect Repair Filing From Warehouse 22 Apr 2016

Process: 7699

Shred Sensitive Paperwork In JL Office 19 May 2016

Process: 7712

Review Inward Payments 01 Jul 2016

Process: 7752

SRS Folder 22 Nov 2016

Process: 7760

Send Service Offers 31 Jan 2017

Process: 7783

PDF VST Invoices And Purchase Orders 10 Feb 2017

Process: 7792

Shipped Order Success Report 13 Mar 2017

Process: 7795

Answering UK Web Questions 27 Apr 2017

Process: 5859

Review Un-shipped Parcels 17 Feb 2016

Process: 6954

Back Orders Review - By Customer 09 Mar 2016

Process: 7748

Check Repair Orders 10 Oct 2016

Process: 7749

Check Repair Quotes 10 Oct 2016

Process: 7838

Review VIAMED Feedback - Customer Feedback Negative 23 Sep 2017

Process: 7839

Review VIAMED Feedback - Customer Complaints 23 Sep 2017

Process: 7840

Review VST Feedback - Customer Feedback Negative 23 Sep 2017

Process: 7841

Review VST Feedback - Customer Complaints 23 Sep 2017

Process: 7842

Review VIAMED Product Feedback Negative 23 Sep 2017

Process: 7843

Review VST Product Feedback Negative 23 Sep 2017

		Process: 7872 Embargo Countries NOT Allowed To Sell To 16 Oct 2017
5.2 Policy		
5.2.1 Top management shall establish, implement and maintain a quality policy that: a) is appropriate to the purpose and context of the organization and supports its strategic direction; b) provides a framework for setting quality objectives; c) includes a commitment to satisfy applicable requirements; d) includes a commitment to continual improvement of the quality management system. Establishing the quality policy	Top Level Document: VM3COP00.00 Viamed Quality Statement policy and objectives Revision Document ID22684 Date Revision 16 Oct 2017 Reviewed 03 Aug 2021 Audit 20 Process verification to Managment Revision Document ID73324 Date Revision 26 Oct 2021 Reviewed 26 Oct 2021 VM3COP00.01 Company objectives Revision Document ID22842 Date Revision 17 Oct 2017 Reviewed 17 Oct 2017 VM3COP00.00 VST Quality Statement policy and objectives Revision Document ID22062 Date Revision 16 Sep 2017 Reviewed 24 Aug 2021	Process: 7833 Importance Of Effective Quality Management 20 Sep 2017 Process: 7828 Review The Quality Policy Viamed 16 Sep 2017 Process: 7827 Review The Quality Policy VST 16 Sep 2017 Process: 7668 Responsibility Allocation : Upgrading Intrastats ISO Quality system 09 Mar 2016
5.2.2 The quality policy shall: a) be available and be maintained as documented information; b) be communicated, understood and applied within the organization; c) be available to relevant interested parties, as appropriate. Communicating the quality policy	Top Level Document: VOP 01 Documentation and Records, Control, Creation, Storage, Retrieval, Revision Control and Online Records Revision Document ID75407 Date Revision 18 Nov 2021 Reviewed 18 Nov 2021 Audit 10 Documentation Control Revision Document ID63807 Date Revision 30 Jun 2021 Reviewed 30 Jun 2021 VM3COP00.00 VST Quality Statement policy and objectives Revision Document ID22062	Process: 7833 Importance Of Effective Quality Management 20 Sep 2017 Process: 7828 Review The Quality Policy Viamed 16 Sep 2017 Process: 7827 Review The Quality Policy VST 16 Sep 2017 Process: 7676 PDFing Of Invoices Viamed 17 Mar 2016 Process: 7668 Responsibility Allocation : Upgrading Intrastats ISO Quality system 09 Mar 2016

	Date Revision 16 Sep 2017 Reviewed 24 Aug 2021	
5.3 Top management shall ensure that the responsibilities and authorities for relevant roles are assigned, communicated and understood within the organization. Top management shall assign the responsibility and authority for: a) ensuring that the quality management system conforms to the requirements of this International Standard; b) ensuring that the processes are delivering their intended outputs; c) reporting on the performance of the quality management system and on opportunities for improvement (see 10.1), in particular to top management; d) ensuring the promotion of customer focus throughout the organization; e) ensuring that the integrity of the quality management system is maintained when changes to the quality management system are planned and implemented. Organizational roles, responsibilities and authorities	Top Level Document: VOP 02 Personnel and Responsibility , Staff and Staffing Issues, Training, Roles and Tasks Revision Document ID73529 Date Revision 29 Oct 2021 Reviewed 29 Oct 2021 Audit 20 Process verification to Managment Revision Document ID73324 Date Revision 26 Oct 2021 Reviewed 26 Oct 2021	Process: 7744 FDA Device Establishment Registration And Listing 28 Sep 2016 Process: 7740 Weights Per Region Needed To Submit EC Sales List 13 Sep 2016 Process: 7668 Responsibility Allocation : Upgrading Intrastats ISO Quality system 09 Mar 2016 Process: 7387 Responsibility Allocation : VST Stock Meeting Purchase Order Requirements 09 Mar 2016
6 Planning		
6 Planning		Process: 7433 Responsibility Allocation : VST Board Directors Meeting 09 Mar 2016
6.1 Actions to address risks and		

opportunities		
6.1 When planning for the quality management system, the organization shall consider the issues referred to in 4.1 and the requirements referred to in 4.2 and determine the risks and opportunities that need to be addressed to: a) give assurance that the quality management system can achieve its intended result(s); b) enhance desirable effects; c) prevent, or reduce, undesired effects; d) achieve improvement.	Top Level Document: VOP 24 Needs, Risks and Expectations of External Parties Revision Document ID31060 Date Revision 30 Sep 2019 Reviewed 30 Sep 2019 Top Level Document: Need Risks and Expectations of External Parties VST Revision Document ID74925 Date Revision 15 Nov 2021 Reviewed 15 Nov 2021 Audit 18 Management Review Revision Document ID73320 **Date Revision 26 Oct 2021 Reviewed 26 Oct 2021 VM3COP24.01 Definitions of Risk Revision Document ID75525 Date Revision 19 Nov 2021 Reviewed 19 Nov 2021	Process: 7670 Humanmed general Issues 09 Mar 2016
6.1.2 The organization shall plan: a) actions to address these risks and opportunities; b) how to: 1) integrate and implement the actions into its quality management system processes (see 4.4); 2) evaluate the effectiveness of these actions. Actions taken to address risks and opportunities shall be proportionate to the potential impact on the conformity of products and services. NOTE 1 Options to address risks can include avoiding risk, taking risk in order to pursue an opportunity, eliminating the risk source, changing the	Audit 18 Management Review Revision Document ID73320 **Date Revision 26 Oct 2021 Reviewed 26 Oct 2021 Audit 12 CE Files Revision Document ID63815 Date Revision 30 Jun 2021 Reviewed 30 Jun 2021	Process: 7832 Cleardown Emailed Invoices 20 Sep 2017 Process: 7809 Pro-Active Marketing 06 Jun 2017 Process: 7673 Check Expiry Dated Stock 09 Mar 2016 Process: 7664 Responsibility Allocation : Marketing Job Logger 09 Mar 2016 Process: 7394 Responsibility Allocation : VST Stock Meeting Repairs Review - General 09 Mar 2016

likelihood or consequences, sharing the risk, or retaining risk by informed decision. NOTE 2 Opportunities can lead to the adoption of new practices, launching new products, opening new markets, addressing new customers, building partnerships, using new technology and other desirable and viable possibilities to address the organization's or its customers' needs.		
6.2 Quality objectives and planning to achieve them		
6.2.1 The organization shall establish quality objectives at relevant functions, levels and processes needed for the quality management system. The quality objectives shall: a) be consistent with the quality policy; b) be measurable; c) take into account applicable requirements; d) be relevant to conformity of products and services and to enhancement of customer satisfaction; e) be monitored; f) be communicated; g) be updated as appropriate. The organization shall maintain documented information on the quality objectives	Top Level Document: VOP 13 Process Monitoring, System Reviews, Audits, Management Reviews and Analysis Data Revision Document ID75461 Date Revision 18 Nov 2021 Reviewed 18 Nov 2021 Audit 10 Documentation Control Revision Document ID63807 Date Revision 30 Jun 2021 Reviewed 30 Jun 2021 Audit 20 Process verification to Managment Revision Document ID73324 Date Revision 26 Oct 2021 Reviewed 26 Oct 2021	Process: 7830 Review Q.A. Failures Report 18 Sep 2017 Process: 7828 Review The Quality Policy Viamed 16 Sep 2017 Process: 7827 Review The Quality Policy VST 16 Sep 2017 Process: 7825 Responsibility Allocation : Order Picking 06 Sep 2017 Process: 7822 Review Oxylink Stock 26 Jul 2017 Process: 7797 Check Order Are Being Picked In Priority Order 10 May 2017 Process: 7761 Send VST Delivery Notifications 01 Feb 2017 Process: 7760 Send Service Offers 31 Jan 2017 Process: 7734 Responsibility Allocation : Humanmed Order Processing 25 Aug 2016 Process: 7710 Responsibility Allocation : Proforma And Quote Processing 29 Jun 2016 Process: 7709

Delivered not Invoiced 28 Jun 2016

Process: 7953

Vandagraph Delivery Notifications 26 May 2020

Process: 7691

Ship Sale Or Returns 21 Apr 2016

Process: 7690

Ship Repairs 21 Apr 2016

Process: 7686

Thorough Checking Of Awaiting Action Tray 21 Apr 2016

Process: 7685

Repairs Ready For Invoice 18 Apr 2016

Process: 7684

Repairs Ready For Quote 18 Apr 2016

Process: 7683

Check Stock For Proforma 18 Apr 2016

Process: 7678

Check Catalog 360 Circle For Quotes And Orders 08 Apr 2016

Process: 7674

Check Repairs Ready For Invoice List 10 Mar 2016

Process: 7673

Check Expiry Dated Stock 09 Mar 2016

Process: 7670

Humanmed general Issues 09 Mar 2016

Process: 7668

Responsibility Allocation : Upgrading Intrastats ISO Quality system
09 Mar 2016

Process: 7398

Responsibility Allocation : VST Stock Meeting UPS Shipping Fuel
Surcharge 09 Mar 2016

Process: 7396

Responsibility Allocation : VST Stock Meeting `Goods Out` Review
09 Mar 2016

Process: 7394

Responsibility Allocation : VST Stock Meeting Repairs Review -
General 09 Mar 2016

Process: 7389

Responsibility Allocation : VST Stock Meeting Returns Overview -
From Customers 09 Mar 2016

Process: 7387

Responsibility Allocation : VST Stock Meeting Purchase Order Requirements 09 Mar 2016
Process: 7385
Responsibility Allocation : VST Stock Meeting Sales Forward Orders Review 09 Mar 2016
Process: 6938
Responsibility Allocation : Customer Database Updates 09 Mar 2016
Process: 6956
Responsibility Allocation : Sales Order Issues 09 Mar 2016
Process: 7090
Responsibility Allocation : Office Procedures 09 Mar 2016
Process: 6898
GHX Web Pricing 09 Mar 2016
Process: 5871
Check Sale Or Returns 17 Feb 2016
Process: 5876
E.Commerce Cardea And Multiquote 17 Feb 2016
Process: 5872
Check Sale Or Returns Export 17 Feb 2016
Process: 2
Answering Telephones 16 Feb 2016
Process: 3
Responsibility Allocation : Meeting And Greeting Visitors To The Company 16 Feb 2016
Process: 4
Responsibility Allocation : Assisting With Refreshments For Visitors 16 Feb 2016
Process: 5
Responsibility Allocation : Processing Of Sales Orders 16 Feb 2016
Process: 6
Responsibility Allocation : Updating Contact Management System 16 Feb 2016
Process: 7
Responsibility Allocation : Checking Of Sales Orders 16 Feb 2016
Process: 8
Responsibility Allocation : Order And Status Liaison With Customers 16 Feb 2016
Process: 10
Distribution Of Emails 16 Feb 2016

Process: 11
Distribution Of Mail 16 Feb 2016

Process: 14
Fax Paper 16 Feb 2016

Process: 15
Filing and Archiving 16 Feb 2016

Process: 16
Responsibility Allocation : Photocopying 16 Feb 2016

Process: 21
Office Sales Projects 16 Feb 2016

Process: 36
Emailing Of Invoices 16 Feb 2016

Process: 5875
Check Paypal For Orders 17 Feb 2016

Process: 5879
Responsibility Allocation : Customer Returning Goods On Our UPS Account 18 Feb 2016

Process: 5882
Responsibility Allocation : Send Post To Humanmed 24 Feb 2016

Process: 5891
Processing Of Repair Quotes And Orders 25 Feb 2016

Process: 5892
Checking EBay And Amazon For Orders And Messages 25 Feb 2016

Process: 5893
Answering Website Questions 25 Feb 2016

Process: 5894
Checking Of Active List 25 Feb 2016

Process: 5895
Responsibility Allocation : Completing Office Job List 25 Feb 2016

Process: 5896
Responsibility Allocation : Ensuring ORD's Are Taken To Goods Out And Invoices Are Retrieved 25 Feb 2016

Process: 5899
Proforma And Quote Chasing 25 Feb 2016

Process: 5901
Link Call Log Contacts To The CRM 02 Mar 2016

Process: 5913
Check For Humanmed Orders In Logistics Mailbox 03 Mar 2016

Process: 5943

Check Cardea And Multiquote 08 Mar 2016
Process: 5944
Responsibility Allocation : Chasing Lost Customers 08 Mar 2016
Process: 5945
Responsibility Allocation : Sending Samples 08 Mar 2016
Process: 5946
Responsibility Allocation : Sending Sale Or Returns 08 Mar 2016
Process: 5947
Responsibility Allocation : Search For Distributors 08 Mar 2016
Process: 5948
Adding New Accounts To Opera 08 Mar 2016
Process: 5949
Filling Credit Card Slips 08 Mar 2016
Process: 6958
Responsibility Allocation : Shipped Order Queries 09 Mar 2016
Process: 7676
PDFing Of Invoices Viamed 17 Mar 2016
Process: 7693
Collect Repair Filing From Warehouse 22 Apr 2016
Process: 7699
Shred Sensitive Paperwork In JL Office 19 May 2016
Process: 7712
Review Inward Payments 01 Jul 2016
Process: 7735
Ensure SOR's Are Followed Up 01 Sep 2016
Process: 7752
SRS Folder 22 Nov 2016
Process: 7758
Check For GHX Orders 17 Jan 2017
Process: 7783
PDF VST Invoices And Purchase Orders 10 Feb 2017
Process: 7795
Answering UK Web Questions 27 Apr 2017
Process: 5859
Review Un-shipped Parcels 17 Feb 2016
Process: 6954
Back Orders Review - By Customer 09 Mar 2016
Process: 7748
Check Repair Orders 10 Oct 2016

		Process: 7749 Check Repair Quotes 10 Oct 2016
6.2.2 When planning how to achieve its quality objectives, the organization shall determine: a) what will be done; b) what resources will be required; c) who will be responsible; d) when it will be completed; e) how the results will be evaluated.	Audit 20 Process verification to Managment Revision Document ID73324 Date Revision 26 Oct 2021 Reviewed 26 Oct 2021	Process: 7387 Responsibility Allocation : VST Stock Meeting Purchase Order Requirements 09 Mar 2016
6.3 When the organization determines the need for changes to the quality management system, the changes shall be carried out in a planned manner (see 4.4). The organization shall consider: a) the purpose of the changes and their potential consequences; b) the integrity of the quality management system; c) the availability of resources; d) the allocation or reallocation of responsibilities and authorities. Planning of changes	Top Level Document: VOP 02 Personnel and Responsibility , Staff and Staffing Issues, Training, Roles and Tasks Revision Document ID73529 Date Revision 29 Oct 2021 Reviewed 29 Oct 2021 Top Level Document: VOP 01 Documentation and Records, Control, Creation, Storage, Retrieval, Revision Control and Online Records Revision Document ID75407 Date Revision 18 Nov 2021 Reviewed 18 Nov 2021 Audit 10 Documentation Control Revision Document ID63807 Date Revision 30 Jun 2021 Reviewed 30 Jun 2021 Upgrading of the ISO Systems 2016 - 2017 Revision Document ID22140 Date Revision 20 Sep 2017 Reviewed 20 Sep 2017 Explanation Employee Roles Titles Responsibilitys Processes and Repeating Tasks Monitoring Revision Document ID22287	

7 Support

7 Support		
7.1 Resources		
7.1.1 General The organization shall determine and provide the resources needed for the establishment, implementation, maintenance and continual improvement of the quality management system. The organization shall consider: a) the capabilities of, and constraints on, existing internal resources; b) what needs to be obtained from external providers. General	Audit 18 Management Review Revision Document ID73320 **Date Revision 26 Oct 2021 Reviewed 26 Oct 2021	Process: 7814 Responsibility Allocation : Viamed Repairs 06 Jun 2017 Process: 7670 Humanmed general Issues 09 Mar 2016
7.1.2 The organization shall determine and provide the persons necessary for the effective implementation of its quality management system and for the operation and control of its processes. People	Top Level Document: VOP 12 Training Revision Document ID31024 Date Revision 30 Sep 2019 Reviewed 30 Sep 2019 Audit 08 Training, Competence and Human Resources Revision Document ID70147 Date Revision 20 Sep 2021 Reviewed 20 Sep 2021 Audit 20 Process verification to Managment Revision Document ID73324 Date Revision 26 Oct 2021 Reviewed 26 Oct 2021 Employee Roles Revision Document ID20125	Process: 7713 Review Roles And Responsibilitys 17 Aug 2016 Process: 7793 Team Review Meeting 16 Mar 2017 Process: 7759 Health Declaration Sheet 23 Jan 2017 Process: 7670 Humanmed general Issues 09 Mar 2016

	Date Revision 16 May 2017 Reviewed 16 May 2017	
7.1.3 The organization shall determine, provide and maintain the infrastructure necessary for the operation of its processes and to achieve conformity of products and services. NOTE Infrastructure can include: a) buildings and associated utilities; b) equipment, including hardware and software; c) transportation resources; d) information and communication technology. Infrastructure	Top Level Document: VOP 11 Equipment Control, Office, Warehouse, Pcs and Equipment Revision Document ID31008 Date Revision 30 Sep 2019 Reviewed 30 Sep 2019 Top Level Document: VOP 18 Maintenance Building, Fabric and Infrastructure Revision Document ID31036 Date Revision 30 Sep 2019 Reviewed 30 Sep 2019 Top Level Document: VOP 06 Measurement Control Viamed VST, Calibration, QA Stock Revision Document ID53615 Date Revision 11 Feb 2021 Reviewed 11 Feb 2021 Audit 10 Documentation Control Revision Document ID63807 Date Revision 30 Jun 2021 Reviewed 30 Jun 2021 Employee Roles Revision Document ID20125 Date Revision 16 May 2017 Reviewed 16 May 2017 Ghyll House Fire Certificate Revision Document ID12303 Date Revision 15 Mar 2013 Reviewed 15 Mar 2013 HSE Fire / Exit Escape route Basement floor plans Revision Document ID15401 Date Revision 07 Aug 2015 Reviewed 28 Sep 2020 HSE Fire / Exit Escape route Ghyll House floor plans	Process: 7091 Calibration Index 09 Mar 2016 Process: 7745 UPS Invoices Viamed 06 Oct 2016 Process: 7746 UPS Invoices VST 06 Oct 2016 Process: 7747 UPS Invoices Vandagraph 06 Oct 2016 Process: 7120 General Maintenance Requirements 09 Mar 2016 Process: 5940 Thumb Nail Processor 07 Mar 2016 Process: 7739 Intrastats Amendment Log 12 Sep 2016 Process: 7129 Intrastats Cross Reference Database Tables Updates 09 Mar 2016 Process: 7126 Intrastats Requested Page updates 09 Mar 2016 Process: 5905 Responsibility Allocation : Price Checking 02 Mar 2016 Process: 5866 UPS Shipping Fuel Surcharge 17 Feb 2016 Process: 6972 UPS Shipping Fuel Surcharge 09 Mar 2016 Process: 5903 Responsibility Allocation : Weather Station 02 Mar 2016 Process: 7711 Import Bank CSV 01 Jul 2016 Process: 7706 Update Virus Software And Scan For Viruses 10 Jun 2016 Process: 46 Responsibility Allocation : Backup Server Status 16 Feb 2016 Process: 48 Responsibility Allocation : Internet 16 Feb 2016 Process: 45 Responsibility Allocation : Main Server Status 16 Feb 2016 Process: 44

Revision Document ID27948
Date Revision 29 Oct 2018 Reviewed 29 Oct 2018
VM3COP20.35 Ups Calculator
Revision Document ID17149
Date Revision 05 Jul 2016 Reviewed 05 Jul 2016
VM3COP03.05 Procedures for customer returning goods on our UPS account number
Revision Document ID17155
Date Revision 05 Jul 2016 Reviewed 05 Jul 2016
Audit 15 Production
Revision Document ID59614
Date Revision 11 May 2021 Reviewed 11 May 2021
FIRE Report Premisis
Revision Document ID61402
Date Revision 02 Jun 2021 Reviewed 02 Jun 2021
HSE Fire appliances HSE Fire Exit / Escape Route Ground Floor plans
Revision Document ID27944
Date Revision 29 Oct 2018 Reviewed 29 Oct 2018
HSE Fire Risk Assessment
Revision Document ID21790
Date Revision 04 Sep 2017 Reviewed 04 Sep 2017
Audit 19 Health and Safety, Working Conditions and Building Fabric Issues
Revision Document ID68045
Date Revision 24 Aug 2021 Reviewed 24 Aug 2021
CPM 21 Fire Exit / Escape Route Procedures
Revision Document ID21892
Date Revision 07 Sep 2017 Reviewed 07

Secure Socket Level Certificate 16 Feb 2016
Process: 49
Responsibility Allocation : Wifi 16 Feb 2016
Process: 50
Responsibility Allocation : Guest Access Wifi 16 Feb 2016
Process: 5941
Responsibility Allocation : Replace Main Server 07 Mar 2016
Process: 5939
Responsibility Allocation : Email ISP Routing 05 Mar 2016
Process: 7121
Responsibility Allocation : General Computer Maintenance 09 Mar 2016
Process: 7125
Responsibility Allocation : Intrastats Urgent Problems 09 Mar 2016
Process: 7124
Responsibility Allocation : Intrastats 09 Mar 2016
Process: 7127
Responsibility Allocation : Intrastats Unfinished in progress Processes 09 Mar 2016
Process: 7128
Responsibility Allocation : Intrastats Future Features needed 09 Mar 2016
Process: 7133
Responsibility Allocation : Intrastats Contact Manager 09 Mar 2016
Process: 7704
Responsibility Allocation : Computer Failure Diagnostics 24 May 2016
Process: 7835
Electrics Need Checking 20 Sep 2017
Process: 7836
Central Heating For Winter 20 Sep 2017
Process: 7832
Cleardown Emailed Invoices 20 Sep 2017
Process: 7823
Saftey Tester Data 02 Aug 2017
Process: 7805
Empty Kitchen Bins 22 May 2017
Process: 7804
Sweep Kitchen Floor 22 May 2017

Sep 2017

Explanation Employee Roles and Titles

Revision Document ID22144

Date Revision 20 Sep 2017 Reviewed 20 Sep 2017

HSE Fire Exit / Escape Route Ground Floor plans Document

Revision Document ID2558

Date Revision 01 Aug 2007 Reviewed 01 Aug 2007

DO NOT USE VM3COP11 Calibration

Revision Document ID8713

Date Revision 12 Oct 2011 Reviewed 12 Oct 2011

VM3COP20.07 UPS Procedures

Revision Document ID8722

Date Revision 12 Oct 2011 Reviewed 12 Oct 2011

HSE Fire Safety Risk Assessment

Revision Document ID892

Date Revision 25 Oct 2006 Reviewed 25 Oct 2006

Process: 7803

Dishwashing 22 May 2017

Process: 7802

Clean Kitchen Sides 22 May 2017

Process: 7756

Carbon Monoxide Alarm 05 Jan 2017

Process: 7742

Boiler Check 26 Sep 2016

Process: 7698

Clean Toilets 17 May 2016

Process: 7687

Vandagraph Duckets 21 Apr 2016

Process: 7672

Off Site Backup 09 Mar 2016

Process: 7402

Responsibility Allocation : VST Calibration P.A.T. Testing 09 Mar 2016

Process: 7401

Responsibility Allocation : VST Calibration 09 Mar 2016

Process: 7857

Software Validation Stock Tracking Check 01 Oct 2017

Process: 5851

Duplicate Documents 17 Feb 2016

Process: 59

Out Of Date Documents 17 Feb 2016

Process: 7850

Software Validation Scan In Correct Product 01 Oct 2017

Process: 7851

****Software Validation Scan Un-QA Product To Order 23 Nov 2021**

Process: 7852

Software Validation Expired Stock 01 Oct 2017

Process: 7853

Software Validation Non Sell Able Shelf 01 Oct 2017

Process: 7854

Software Validation In Production List 01 Oct 2017

Process: 7855

Software Validation - Production Lists 01 Oct 2017

Process: 7856

Software Validation Unchecked Orders 01 Oct 2017

		Process: 7870 Software Validation Non Conformance Product Risk Feedback Loop 15 Oct 2017 Process: 7869 Hand Drill Checklist 13 Oct 2017 Process: 7868 Pillar Drill Checklist 13 Oct 2017 Process: 7867 Bandsaw Checklist 13 Oct 2017 Process: 7866 Oxygen Cylinder Check 13 Oct 2017 Process: 7865 Software Validation Conflicting Audits 07 Oct 2017 Process: 7864 ESD Work Stations 07 Oct 2017
7.1.4 The organization shall determine, provide and maintain the environment necessary for the operation of its processes and to achieve conformity of products and services. NOTE A suitable environment can be a combination of human and physical factors, such as: a) social (e.g. non-discriminatory, calm, non-confrontational); b) psychological (e.g. stress-reducing, burnout prevention, emotionally protective); c) physical (e.g. temperature, heat, humidity, light, airflow, hygiene, noise). These factors can differ substantially depending on the products and services provided. Environment for the operation of processes	Top Level Document: VOP 12 Training Revision Document ID31024 Date Revision 30 Sep 2019 Reviewed 30 Sep 2019 Top Level Document: VOP 16 Health and Safety, Company Personnel Manual Revision Document ID31032 Date Revision 30 Sep 2019 Reviewed 30 Sep 2019 Top Level Document: VOP 18 Maintenance Building, Fabric and Infrastructure Revision Document ID31036 Date Revision 30 Sep 2019 Reviewed 30 Sep 2019 Audit 19 Health and Safety, Working Conditions and Building Fabric Issues Revision Document ID68045 Date Revision 24 Aug 2021 Reviewed 24 Aug 2021 Audit 08 Training, Competence and Human Resources Revision Document ID70147 Date Revision 20 Sep 2021 Reviewed 20	Process: 7750 Meeting With Management 14 Oct 2016 Process: 7120 General Maintenance Requirements 09 Mar 2016 Process: 7753 Management Meeting Warehouse 22 Nov 2016 Process: 7836 Central Heating For Winter 20 Sep 2017 Process: 7811 Responsibility Allocation : General Area 06 Jun 2017 Process: 7806 Watering Plants 22 May 2017 Process: 7698 Clean Toilets 17 May 2016 Process: 7845 7.1.4 Environment Of Operations 25 Sep 2017

Sep 2021
Fire risk assessment 15/17 Station Road
 Revision Document ID48624
 Date Revision 26 Nov 2020 Reviewed 26 Nov 2020
CPM 25 Health and Safety Policy
Viamed
 Revision Document ID14332
 Date Revision 25 Sep 2014 Reviewed 04 Sep 2017
Audit 07 Handling and Storage
 Revision Document ID58347
 Date Revision 23 Apr 2021 Reviewed 23 Apr 2021
Audit 19 Health and Safety, Working Conditions and Building Fabric Issues
 Revision Document ID68045
 Date Revision 24 Aug 2021 Reviewed 24 Aug 2021
CPM 39 Smoking Policy
 Revision Document ID6782
 Date Revision 15 Feb 2010 Reviewed 15 Feb 2010
CPM 16 Dress Code
 Revision Document ID7055
 Date Revision 26 Apr 2010 Reviewed 22 Jul 2014
CPM 15 Disciplinary Procedures
 Revision Document ID25502
 Date Revision 05 Mar 2018 Reviewed 05 Mar 2018
Audit 08 Training, Competence and Human Resources
 Revision Document ID70147
 Date Revision 20 Sep 2021 Reviewed 20 Sep 2021

7.1.5
Monitoring and measuring resources

7.1.5.1 7.1.5.1 General The organization shall determine and provide the resources needed to ensure valid and reliable results when monitoring or measuring is used to verify the conformity of products and services to requirements. The organization shall ensure that the resources provided: a) are suitable for the specific type of monitoring and measurement activities being undertaken; b) are maintained to ensure their continuing fitness for their purpose. The organization shall retain appropriate documented information as evidence of fitness for purpose of the monitoring and measurement resources. General	Top Level Document: VOP 06 Measurement Control Viamed VST, Calibration, QA Stock Revision Document ID53615 Date Revision 11 Feb 2021 Reviewed 11 Feb 2021 Audit 06 Calibration Revision Document ID63048 Date Revision 22 Jun 2021 Reviewed 22 Jun 2021 Audit 07 Handling and Storage Revision Document ID58347 Date Revision 23 Apr 2021 Reviewed 23 Apr 2021	Process: 6949 Responsibility Allocation : VIAMED Stock Meeting QA Processing 09 Mar 2016 Process: 7689 Move Stock From QA Shelf To Stock Shelf Monday 21 Apr 2016 Process: 7694 Move Stock From QA Shelf To Stock Shelf Tuesday 28 Apr 2016 Process: 7695 Top Up Quick Shipping Shelves 28 Apr 2016 Process: 7830 Review Q.A. Failures Report 18 Sep 2017 Process: 7794 V1000 Commissions Review 30 Mar 2017 Process: 7705 Checking For Uploaded Files 08 Jun 2016 Process: 7690 Ship Repairs 21 Apr 2016 Process: 7676 PDFing Of Invoices Viamed 17 Mar 2016 Process: 7673 Check Expiry Dated Stock 09 Mar 2016 Process: 7670 Humanmed general Issues 09 Mar 2016 Process: 7394 Responsibility Allocation : VST Stock Meeting Repairs Review - General 09 Mar 2016
7.1.5.2 When measurement traceability is a requirement, or is considered by the organization to be an essential part of providing confidence in the validity of measurement results, measuring equipment shall be: a) calibrated or verified, or both, at specified intervals, or prior to use, against measurement standards traceable to international or national measurement standards; when no such standards exist, the	Top Level Document: VOP 06 Measurement Control Viamed VST, Calibration, QA Stock Revision Document ID53615 Date Revision 11 Feb 2021 Reviewed 11 Feb 2021 Top Level Document: VOP 07 Stock Control, Handling, Control of Labelling, Storage, Movement Revision Document ID75973 **Date Revision 24 Nov 2021 Reviewed 24 Nov 2021 Top Level Document: VOP 11	Process: 7830 Review Q.A. Failures Report 18 Sep 2017 Process: 7823 Safety Tester Data 02 Aug 2017 Process: 7814 Responsibility Allocation : Viamed Repairs 06 Jun 2017 Process: 7813 Responsibility Allocation : VST Repairs 06 Jun 2017 Process: 7812 Responsibility Allocation : Vandagraph Repairs 06 Jun 2017 Process: 7798 Orders And Items Shipped Per Month 10 May 2017 Process: 7744

basis used for calibration or verification shall be retained as documented information; b) identified in order to determine their status; c) safeguarded from adjustments, damage or deterioration that would invalidate the calibration status and subsequent measurement results. The organization shall determine if the validity of previous measurement results has been adversely affected when measuring equipment is found to be unfit for its intended purpose, and shall take appropriate action as necessary Measurement traceability	Equipment Control, Office, Warehouse, Pcs and Equipment Revision Document ID31008 Date Revision 30 Sep 2019 Reviewed 30 Sep 2019 Audit 06 Calibration Revision Document ID63048 Date Revision 22 Jun 2021 Reviewed 22 Jun 2021 Audit 10 Documentation Control Revision Document ID63807 Date Revision 30 Jun 2021 Reviewed 30 Jun 2021	FDA Device Establishment Registration And Listing 28 Sep 2016 Process: 7705 Checking For Uploaded Files 08 Jun 2016 Process: 7693 Collect Repair Filing From Warehouse 22 Apr 2016 Process: 7692 Responsibility Allocation : Take Complete Repair Paperwork To Office 22 Apr 2016 Process: 7673 Check Expiry Dated Stock 09 Mar 2016 Process: 7670 Humanmed general Issues 09 Mar 2016 Process: 7401 Responsibility Allocation : VST Calibration 09 Mar 2016 Process: 7048 Control of monitoring and measuring devices 09 Mar 2016
7.1.6 The organization shall determine the knowledge necessary for the operation of its processes and to achieve conformity of products and services. This knowledge shall be maintained and be made available to the extent necessary. When addressing changing needs and trends, the organization shall consider its current knowledge and determine how to acquire or access any necessary additional knowledge and required updates. NOTE 1 Organizational knowledge is knowledge specific to the organization; it is generally gained by experience. It is information that is used and shared to achieve the organization's objectives. NOTE 2 Organizational knowledge can be based on:	Audit 08 Training, Competence and Human Resources Revision Document ID70147 Date Revision 20 Sep 2021 Reviewed 20 Sep 2021 Audit 10 Documentation Control Revision Document ID63807 Date Revision 30 Jun 2021 Reviewed 30 Jun 2021 Audit 12 CE Files Revision Document ID63815 Date Revision 30 Jun 2021 Reviewed 30 Jun 2021	Process: 7830 Review Q.A. Failures Report 18 Sep 2017 Process: 7744 FDA Device Establishment Registration And Listing 28 Sep 2016 Process: 7673 Check Expiry Dated Stock 09 Mar 2016 Process: 7670 Humanmed general Issues 09 Mar 2016 Process: 7387 Responsibility Allocation : VST Stock Meeting Purchase Order Requirements 09 Mar 2016 Process: 7863 Maintain Repair Codes List 05 Oct 2017

a) internal sources (e.g. intellectual property; knowledge gained from experience; lessons learned from failures and successful projects; capturing and sharing undocumented knowledge and experience; the results of improvements in processes, products and services); b) external sources (e.g. standards; academia; conferences; gathering knowledge from customers or external providers) Organizational knowledge		
7.2 7.2 Competence The organization shall: a) determine the necessary competence of person(s) doing work under its control that affects the performance and effectiveness of the quality management system; b) ensure that these persons are competent on the basis of appropriate education, training, or experience; c) where applicable, take actions to acquire the necessary competence, and evaluate the effectiveness of the actions taken; d) retain appropriate documented information as evidence of competence. NOTE Applicable actions can include, for example, the provision of training to, the mentoring of, or the reassignment of currently employed persons; or the hiring or contracting of competent persons. Competence	Top Level Document: VOP 12 Training Revision Document ID31024 Date Revision 30 Sep 2019 Reviewed 30 Sep 2019 Top Level Document: VOP 02 Personnel and Responsibility , Staff and Staffing Issues, Training, Roles and Tasks Revision Document ID73529 Date Revision 29 Oct 2021 Reviewed 29 Oct 2021 Audit 08 Training, Competence and Human Resources Revision Document ID70147 Date Revision 20 Sep 2021 Reviewed 20 Sep 2021 Audit 19 Health and Safety, Working Conditions and Building Fabric Issues Revision Document ID68045 Date Revision 24 Aug 2021 Reviewed 24 Aug 2021 Explanation Employee Roles and Titles Revision Document ID22144 Date Revision 20 Sep 2017 Reviewed 20 Sep 2017 Audit 08 Training, Competence and Human Resources	Process: 7673 Check Expiry Dated Stock 09 Mar 2016

	Revision Document ID70147 Date Revision 20 Sep 2021 Reviewed 20 Sep 2021	
7.3 The organization shall ensure that persons doing work under the organization's control are aware of: a) the quality policy; b) relevant quality objectives; c) their contribution to the effectiveness of the quality management system, including the benefits of improved performance; d) the implications of not conforming with the quality management system requirements. Awareness	Top Level Document: VOP 12 Training Revision Document ID31024 Date Revision 30 Sep 2019 Reviewed 30 Sep 2019 Top Level Document: VOP 02 Personnel and Responsibility , Staff and Staffing Issues, Training, Roles and Tasks Revision Document ID73529 Date Revision 29 Oct 2021 Reviewed 29 Oct 2021 Audit 19 Health and Safety, Working Conditions and Building Fabric Issues Revision Document ID68045 Date Revision 24 Aug 2021 Reviewed 24 Aug 2021 Explanation Employee Roles and Titles Revision Document ID22144 Date Revision 20 Sep 2017 Reviewed 20 Sep 2017 Audit 08 Training, Competence and Human Resources Revision Document ID70147 Date Revision 20 Sep 2021 Reviewed 20 Sep 2021	Process: 7673 Check Expiry Dated Stock 09 Mar 2016 Process: 7668 Responsibility Allocation : Upgrading Intrastats ISO Quality system 09 Mar 2016
7.4 7.4 Communication The organization shall determine the internal and external communications relevant to the quality management system, including: a) on what it will communicate; b) when to communicate; c) with whom to communicate; d) how to communicate; e) who communicates. Communication	Audit 10 Documentation Control Revision Document ID63807 Date Revision 30 Jun 2021 Reviewed 30 Jun 2021 Audit 08 Training, Competence and Human Resources Revision Document ID70147 Date Revision 20 Sep 2021 Reviewed 20 Sep 2021 VM3COP27.01 Searching Intrastats Issues	Process: 7673 Check Expiry Dated Stock 09 Mar 2016

	Revision Document ID6657 Date Revision 02 Nov 2009 Reviewed 02 Nov 2009 VM3COP27.17 Complete Auto_calender Issues Revision Document ID16995 Date Revision 26 May 2016 Reviewed 26 May 2016 VM3COP27.36 Auto Close Issues Revision Document ID17082 Date Revision 24 Jun 2016 Reviewed 24 Jun 2016 Overview Issues Meeting Headers List Revision Document ID22169 Date Revision 22 Sep 2017 Reviewed 22 Sep 2017 Issues Overview Revision Document ID23112 Date Revision 22 Oct 2017 Reviewed 22 Oct 2017	
7.5 Documented information		
7.5.1 7.5.1 General The organization's quality management system shall include: a) documented information required by this International Standard; b) documented information determined by the organization as being necessary for the effectiveness of the quality management system. NOTE The extent of documented information for a quality management system can differ from one organization to another due to: ◆◆◆◆the size of organization and its type of activities, processes, products and	Top Level Document: VOP 01 Documentation and Records, Control, Creation, Storage, Retrieval, Revision Control and Online Records Revision Document ID75407 Date Revision 18 Nov 2021 Reviewed 18 Nov 2021 Top Level Document: VM3COP00.00 Viamed Quality Statement policy and objectives Revision Document ID22684 Date Revision 16 Oct 2017 Reviewed 03 Aug 2021 Audit 10 Documentation Control Revision Document ID63807 Date Revision 30 Jun 2021 Reviewed 30	Process: 7744 FDA Device Establishment Registration And Listing 28 Sep 2016 Process: 7734 Responsibility Allocation : Humanmed Order Processing 25 Aug 2016 Process: 7710 Responsibility Allocation : Proforma And Quote Processing 29 Jun 2016 Process: 7709 Delivered not Invoiced 28 Jun 2016 Process: 7953 Vandagraph Delivery Notifications 26 May 2020 Process: 7693 Collect Repair Filing From Warehouse 22 Apr 2016 Process: 7692 Responsibility Allocation : Take Complete Repair Paperwork To

services;

◆◆◆◆the complexity of processes and their interactions;

◆ the competence of persons. **General**

Jun 2021

Audit 20 Process verification to Managment

Revision Document ID73324

Date Revision 26 Oct 2021 Reviewed 26 Oct 2021

Audit 10 Documentation Control

Revision Document ID63807

Date Revision 30 Jun 2021 Reviewed 30 Jun 2021

Audit 20 Process verification to Managment

Revision Document ID73324

Date Revision 26 Oct 2021 Reviewed 26 Oct 2021

Explanation Quality Objectives

Revision Document ID18483

Date Revision 18 Jan 2017 Reviewed 18 Jan 2017

Audit 20 Process verification to Managment

Revision Document ID73324

Date Revision 26 Oct 2021 Reviewed 26 Oct 2021

VM3COP00.00 VST Quality Statement policy and objectives

Revision Document ID22062

Date Revision 16 Sep 2017 Reviewed 24 Aug 2021

Explanation Employee Roles and Titles

Revision Document ID22144

Date Revision 20 Sep 2017 Reviewed 20 Sep 2017

VM3COP00.01 Company objectives

Revision Document ID22842

Date Revision 17 Oct 2017 Reviewed 17 Oct 2017

Office 22 Apr 2016

Process: 7690

Ship Repairs 21 Apr 2016

Process: 7686

Thorough Checking Of Awaiting Action Tray 21 Apr 2016

Process: 7685

Repairs Ready For Invoice 18 Apr 2016

Process: 7684

Repairs Ready For Quote 18 Apr 2016

Process: 7683

Check Stock For Proforma 18 Apr 2016

Process: 7678

Check Catalog 360 Circle For Quotes And Orders 08 Apr 2016

Process: 7674

Check Repairs Ready For Invoice List 10 Mar 2016

Process: 7668

Responsibility Allocation : Upgrading Intrastats ISO Quality system 09 Mar 2016

Process: 7398

Responsibility Allocation : VST Stock Meeting UPS Shipping Fuel Surcharge 09 Mar 2016

Process: 7396

Responsibility Allocation : VST Stock Meeting `Goods Out` Review 09 Mar 2016

Process: 7390

Responsibility Allocation : VST Stock Meeting Returns Overview - Credits 09 Mar 2016

Process: 7385

Responsibility Allocation : VST Stock Meeting Sales Forward Orders Review 09 Mar 2016

Process: 6938

Responsibility Allocation : Customer Database Updates 09 Mar 2016

Process: 6956

Responsibility Allocation : Sales Order Issues 09 Mar 2016

Process: 7090

Responsibility Allocation : Office Procedures 09 Mar 2016

Process: 6898

GHX Web Pricing 09 Mar 2016

Process: 5871

Check Sale Or Returns 17 Feb 2016

Process: 5876

E.Commerce Cardea And Multiquote 17 Feb 2016

Process: 5872

Check Sale Or Returns Export 17 Feb 2016

Process: 2

Answering Telephones 16 Feb 2016

Process: 5

Responsibility Allocation : Processing Of Sales Orders 16 Feb 2016

Process: 6

Responsibility Allocation : Updating Contact Management System 16 Feb 2016

Process: 7

Responsibility Allocation : Checking Of Sales Orders 16 Feb 2016

Process: 8

Responsibility Allocation : Order And Status Liaison With Customers 16 Feb 2016

Process: 9

Distribution Of Faxes 16 Feb 2016

Process: 10

Distribution Of Emails 16 Feb 2016

Process: 11

Distribution Of Mail 16 Feb 2016

Process: 14

Fax Paper 16 Feb 2016

Process: 15

Filing and Archiving 16 Feb 2016

Process: 16

Responsibility Allocation : Photocopying 16 Feb 2016

Process: 21

Office Sales Projects 16 Feb 2016

Process: 36

Emailing Of Invoices 16 Feb 2016

Process: 5875

Check Paypal For Orders 17 Feb 2016

Process: 5879

Responsibility Allocation : Customer Returning Goods On Our UPS Account 18 Feb 2016

Process: 5882

Responsibility Allocation : Send Post To Humanmed 24 Feb 2016
Process: 5891
Processing Of Repair Quotes And Orders 25 Feb 2016
Process: 5892
Checking EBay And Amazon For Orders And Messages 25 Feb 2016
Process: 5893
Answering Website Questions 25 Feb 2016
Process: 5894
Checking Of Active List 25 Feb 2016
Process: 5895
Responsibility Allocation : Completing Office Job List 25 Feb 2016
Process: 5896
Responsibility Allocation : Ensuring ORD's Are Taken To Goods Out
And Invoices Are Retrieved 25 Feb 2016
Process: 5899
Proforma And Quote Chasing 25 Feb 2016
Process: 5901
Link Call Log Contacts To The CRM 02 Mar 2016
Process: 5913
Check For Humanmed Orders In Logistics Mailbox 03 Mar 2016
Process: 5943
Check Cardea And Multiquote 08 Mar 2016
Process: 5944
Responsibility Allocation : Chasing Lost Customers 08 Mar 2016
Process: 5945
Responsibility Allocation : Sending Samples 08 Mar 2016
Process: 5946
Responsibility Allocation : Sending Sale Or Returns 08 Mar 2016
Process: 5947
Responsibility Allocation : Search For Distributors 08 Mar 2016
Process: 5948
Adding New Accounts To Opera 08 Mar 2016
Process: 5949
Filling Credit Card Slips 08 Mar 2016
Process: 6958
Responsibility Allocation : Shipped Order Queries 09 Mar 2016
Process: 7676
PDFing Of Invoices Viamed 17 Mar 2016
Process: 7699

		Shred Sensitive Paperwork In JL Office 19 May 2016 Process: 7712 Review Inward Payments 01 Jul 2016 Process: 7735 Ensure SOR's Are Followed Up 01 Sep 2016 Process: 7752 SRS Folder 22 Nov 2016 Process: 7758 Check For GHX Orders 17 Jan 2017 Process: 7760 Send Service Offers 31 Jan 2017 Process: 7761 Send VST Delivery Notifications 01 Feb 2017 Process: 7783 PDF VST Invoices And Purchase Orders 10 Feb 2017 Process: 7795 Answering UK Web Questions 27 Apr 2017 Process: 7822 Review Oxylink Stock 26 Jul 2017 Process: 5859 Review Un-shipped Parcels 17 Feb 2016 Process: 6954 Back Orders Review - By Customer 09 Mar 2016 Process: 7748 Check Repair Orders 10 Oct 2016 Process: 7749 Check Repair Quotes 10 Oct 2016
7.5.2 7.5.2 Creating and updating When creating and updating documented information, the organization shall ensure appropriate: a) identification and description (e.g. a title, date, author, or reference number); b) format (e.g. language, software version, graphics) and media (e.g. paper, electronic); c) review and approval for suitability and adequacy. Creating and updating	Top Level Document: VOP 01 Documentation and Records, Control, Creation, Storage, Retrieval, Revision Control and Online Records Revision Document ID75407 Date Revision 18 Nov 2021 Reviewed 18 Nov 2021 Top Level Document: VOP 10 Non Conformance, Corrective and Preventive Actions Revision Document ID46915 Date Revision 02 Nov 2020 Reviewed 02	Process: 7782 Remove Started But Not Used Order Numbers 08 Feb 2017 Process: 7676 PDFing Of Invoices Viamed 17 Mar 2016 Process: 7857 Software Validation Stock Tracking Check 01 Oct 2017

	Nov 2020 VM3COP14.01 Disposition of Documents / Records. Revision Document ID15464 Date Revision 14 Aug 2015 Reviewed 14 Aug 2015 Audit 10 Documentation Control Revision Document ID63807 Date Revision 30 Jun 2021 Reviewed 30 Jun 2021 Audit 23 Analysis of Data Revision Document ID67997 Date Revision 23 Aug 2021 Reviewed 23 Aug 2021 DO NOT USE VM3COP01 Document Updates / Amendment control Revision Document ID22201 Date Revision 23 Sep 2017 Reviewed 23 Sep 2017 Guide to Intrastats Revision Document ID24779 Date Revision 22 Dec 2017 Reviewed 22 Dec 2017 Intrastats overview Revision Document ID23567 Date Revision 28 Oct 2017 Reviewed 28 Oct 2017 DO NOT USE VM3COP14 Documentation Revision Document ID9276 Date Revision 18 Oct 2011 Reviewed 18 Oct 2011	
7.5.3 Control of documented information	Top Level Document: VOP 01 Documentation and Records, Control, Creation, Storage, Retrieval, Revision Control and Online Records Revision Document ID75407 Date Revision 18 Nov 2021 Reviewed 18 Nov 2021	Process: 7705 Checking For Uploaded Files 08 Jun 2016

Top Level Document: VOP 10 Non Conformance, Corrective and Preventive Actions
Revision Document ID46915
Date Revision 02 Nov 2020 Reviewed 02 Nov 2020

VM3COP14.01 Disposition of Documents / Records.
Revision Document ID15464
Date Revision 14 Aug 2015 Reviewed 14 Aug 2015

Audit 10 Documentation Control
Revision Document ID63807
Date Revision 30 Jun 2021 Reviewed 30 Jun 2021

Audit 23 Analysis of Data
Revision Document ID67997
Date Revision 23 Aug 2021 Reviewed 23 Aug 2021

DO NOT USE VM3COP01 Document Updates / Amendment control
Revision Document ID22201
Date Revision 23 Sep 2017 Reviewed 23 Sep 2017

Guide to Intrastats
Revision Document ID24779
Date Revision 22 Dec 2017 Reviewed 22 Dec 2017

Intrastats overview
Revision Document ID23567
Date Revision 28 Oct 2017 Reviewed 28 Oct 2017

DO NOT USE VM3COP14 Documentation
Revision Document ID9276
Date Revision 18 Oct 2011 Reviewed 18 Oct 2011

7.5.3.1
Documented information required by the

Audit 10 Documentation Control
Revision Document ID63807

Process: 7744
FDA Device Establishment Registration And Listing 28 Sep 2016

quality management system and by this International Standard shall be controlled to ensure: a) it is available and suitable for use, where and when it is needed; b) it is adequately protected (e.g. from loss of confidentiality, improper use, or loss of integrity).	Date Revision 30 Jun 2021 Reviewed 30 Jun 2021 Audit 20 Process verification to Managment Revision Document ID73324 Date Revision 26 Oct 2021 Reviewed 26 Oct 2021	Process: 7693 Collect Repair Filing From Warehouse 22 Apr 2016 Process: 7692 Responsibility Allocation : Take Complete Repair Paperwork To Office 22 Apr 2016
7.5.3.2 For the control of documented information, the organization shall address the following activities, as applicable: a) distribution, access, retrieval and use; b) storage and preservation, including preservation of legibility; c) control of changes (e.g. version control); d) retention and disposition. Documented information of external origin determined by the organization to be necessary for the planning and operation of the quality management system shall be identified as appropriate, and be controlled. Documented information retained as evidence of conformity shall be protected from unintended alterations. NOTE Access can imply a decision regarding the permission to view the documented information only, or the permission and authority to view and change the documented information.	Audit 10 Documentation Control Revision Document ID63807 Date Revision 30 Jun 2021 Reviewed 30 Jun 2021 Audit 20 Process verification to Managment Revision Document ID73324 Date Revision 26 Oct 2021 Reviewed 26 Oct 2021 Audit 12 CE Files Revision Document ID63815 Date Revision 30 Jun 2021 Reviewed 30 Jun 2021	Process: 7699 Shred Sensitive Paperwork In JL Office 19 May 2016 Process: 7693 Collect Repair Filing From Warehouse 22 Apr 2016 Process: 7692 Responsibility Allocation : Take Complete Repair Paperwork To Office 22 Apr 2016 Process: 7676 PDFing Of Invoices Viamed 17 Mar 2016
8 Operation		
8		Process: 7433

Operation		Responsibility Allocation : VST Board Directors Meeting 09 Mar 2016
8.1 The organization shall plan, implement and control the processes (see 4.4) needed to meet the requirements for the provision of products and services, and to implement the actions determined in Clause 6, by: a) determining the requirements for the products and services; b) establishing criteria for: 1) the processes; 2) the acceptance of products and services; c) determining the resources needed to achieve conformity to the product and service requirements; d) implementing control of the processes in accordance with the criteria; e) determining, maintaining and retaining documented information to the extent necessary: 1) to have confidence that the processes have been carried out as planned; 2) to demonstrate the conformity of products and services to their requirements. The output of this planning shall be suitable for the organizations operations. The organization shall control planned changes and review the consequences of unintended changes, taking action to mitigate any adverse effects, as necessary. The organization shall ensure that outsourced processes are controlled (see 8.4). Operational planning and control	Top Level Document: VOP 08 Production, Reworks, New Production Revision Document ID31072 Date Revision 30 Sep 2019 Reviewed 30 Sep 2019 Top Level Document: VM3COP27.11 Performing a Technical File PMS and risk assessment Revision Document ID75465 Date Revision 18 Nov 2021 Reviewed 18 Nov 2021 VM3COP27.12 Clinical Evaluation Risk assessment Technical Files Revision Document ID15453 Date Revision 11 Aug 2015 Reviewed 11 Aug 2015 Audit 03 Design Control Revision Document ID51631 Date Revision 13 Jan 2021 Reviewed 13 Jan 2021 Audit 07 Handling and Storage Revision Document ID58347 Date Revision 23 Apr 2021 Reviewed 23 Apr 2021 Audit 10 Documentation Control Revision Document ID63807 Date Revision 30 Jun 2021 Reviewed 30 Jun 2021 Audit 23 Analysis of Data Revision Document ID67997 Date Revision 23 Aug 2021 Reviewed 23 Aug 2021 VM3COP24.00 Viamed Overall Risk Analysis Program Revision Document ID47771 Date Revision 12 Nov 2020 Reviewed 12 Nov 2020	Process: 7394 Responsibility Allocation : VST Stock Meeting Repairs Review - General 09 Mar 2016

	Audit 22 Post Market Surveillance Revision Document ID63052 Date Revision 22 Jun 2021 Reviewed 22 Jun 2021	
8.2 Requirements for products and services		Process: 7818 Issues For Accountants - Check Purchasing Journals to see if VAT handled correctly Previous Month 13 Jun 2017 Process: 7819 **Issues For Accountant - Check Contra account 8000 and clear it 08 Dec 2021 Process: 7817 Issues For Accountants - Check suggested invoice report in operas 13 Jun 2017
8.2.1 Communication with customers shall include: a) providing information relating to products and services; b) handling enquiries, contracts or orders, including changes; c) obtaining customer feedback relating to products and services, including customer complaints; d) handling or controlling customer property; e) establishing specific requirements for contingency actions, when relevant. Customer communication	Top Level Document: VOP 03 Contract Review, Enquires, Office Processes Revision Document ID75949 **Date Revision 24 Nov 2021 Reviewed 24 Nov 2021 Top Level Document: VOP 09 Repairs and Servicing Revision Document ID75927 **Date Revision 24 Nov 2021 Reviewed 24 Nov 2021 Top Level Document: VOP 19 FeedBack Customer Complaints Vigilance and Notifications VST Ltd Revision Document ID75995 **Date Revision 24 Nov 2021 Reviewed 24 Nov 2021 Audit 02 Contract Review and Sales Order Processing Revision Document ID69328 Date Revision 09 Sep 2021 Reviewed 09 Sep 2021 Audit 11 Repairs, Servicing and Returns Revision Document ID64142 Date Revision 02 Jul 2021 Reviewed 02 Jul 2021	Process: 7808 Ensure All Invoice Correctly Tagged 02 Jun 2017 Process: 7800 Opera Nominal Ledger Close 11 May 2017 Process: 7790 Humanmed Invoice them For Previous Month 10 Mar 2017 Process: 7789 Withdraw Funds From Paypal 02 Mar 2017 Process: 7783 PDF VST Invoices And Purchase Orders 10 Feb 2017 Process: 7735 Ensure SOR's Are Followed Up 01 Sep 2016 Process: 7734 Responsibility Allocation : Humanmed Order Processing 25 Aug 2016 Process: 7712 Review Inward Payments 01 Jul 2016 Process: 7710 Responsibility Allocation : Proforma And Quote Processing 29 Jun 2016 Process: 7709 Delivered not Invoiced 28 Jun 2016 Process: 7708 Acorn 0014904 17 Jun 2016 Process: 7703

Audit 22 Post Market Surveillance

Revision Document ID63052

Date Revision 22 Jun 2021 Reviewed 22 Jun 2021

VM3COP10.02 Product Recall locate products out in the Field

Revision Document ID74788

Date Revision 12 Nov 2021 Reviewed 12 Nov 2021

Vandagraph Pay Pal Retrieve Funds 23 May 2016

Process: 7702

Responsibility Allocation : Vandagraph Pay Pay Issue Refund 23 May 2016

Process: 7953

Vandagraph Delivery Notifications 26 May 2020

Process: 7691

Ship Sale Or Returns 21 Apr 2016

Process: 7686

Thorough Checking Of Awaiting Action Tray 21 Apr 2016

Process: 7685

Repairs Ready For Invoice 18 Apr 2016

Process: 7684

Repairs Ready For Quote 18 Apr 2016

Process: 7683

Check Stock For Proforma 18 Apr 2016

Process: 7678

Check Catalog 360 Circle For Quotes And Orders 08 Apr 2016

Process: 7674

Check Repairs Ready For Invoice List 10 Mar 2016

Process: 7427

Responsibility Allocation : VST Customer Complaints 09 Mar 2016

Process: 7398

Responsibility Allocation : VST Stock Meeting UPS Shipping Fuel Surcharge 09 Mar 2016

Process: 7396

Responsibility Allocation : VST Stock Meeting `Goods Out` Review 09 Mar 2016

Process: 7391

Responsibility Allocation : VST Stock Meeting Customer Complaints Review **Mandatory** 09 Mar 2016

Process: 7390

Responsibility Allocation : VST Stock Meeting Returns Overview - Credits 09 Mar 2016

Process: 7389

Responsibility Allocation : VST Stock Meeting Returns Overview - From Customers 09 Mar 2016

Process: 7843

Review VST Product Feedback Negative 23 Sep 2017

Process: 7842

Review VIAMED Product Feedback Negative 23 Sep 2017

Process: 7841

Review VST Feedback - Customer Complaints 23 Sep 2017

Process: 7840

Review VST Feedback - Customer Feedback Negative 23 Sep 2017

Process: 7839

Review VIAMED Feedback - Customer Complaints 23 Sep 2017

Process: 7838

Review VIAMED Feedback - Customer Feedback Negative 23 Sep 2017

Process: 7385

Responsibility Allocation : VST Stock Meeting Sales Forward Orders
Review 09 Mar 2016

Process: 6938

Responsibility Allocation : Customer Database Updates 09 Mar 2016

Process: 6956

Responsibility Allocation : Sales Order Issues 09 Mar 2016

Process: 7090

Responsibility Allocation : Office Procedures 09 Mar 2016

Process: 6898

GHX Web Pricing 09 Mar 2016

Process: 5871

Check Sale Or Returns 17 Feb 2016

Process: 5876

E.Commerce Cardea And Multiquote 17 Feb 2016

Process: 5872

Check Sale Or Returns Export 17 Feb 2016

Process: 2

Answering Telephones 16 Feb 2016

Process: 5

Responsibility Allocation : Processing Of Sales Orders 16 Feb 2016

Process: 6

Responsibility Allocation : Updating Contact Management System 16
Feb 2016

Process: 7

Responsibility Allocation : Checking Of Sales Orders 16 Feb 2016

Process: 8

Responsibility Allocation : Order And Status Liaison With Customers

16 Feb 2016

Process: 9

Distribution Of Faxes 16 Feb 2016

Process: 10

Distribution Of Emails 16 Feb 2016

Process: 11

Distribution Of Mail 16 Feb 2016

Process: 14

Fax Paper 16 Feb 2016

Process: 15

Filing and Archiving 16 Feb 2016

Process: 16

Responsibility Allocation : Photocopying 16 Feb 2016

Process: 21

Office Sales Projects 16 Feb 2016

Process: 36

Emailing Of Invoices 16 Feb 2016

Process: 5875

Check Paypal For Orders 17 Feb 2016

Process: 5879

Responsibility Allocation : Customer Returning Goods On Our UPS Account 18 Feb 2016

Process: 5882

Responsibility Allocation : Send Post To Humanmed 24 Feb 2016

Process: 5891

Processing Of Repair Quotes And Orders 25 Feb 2016

Process: 5892

Checking EBay And Amazon For Orders And Messages 25 Feb 2016

Process: 5893

Answering Website Questions 25 Feb 2016

Process: 5894

Checking Of Active List 25 Feb 2016

Process: 5895

Responsibility Allocation : Completing Office Job List 25 Feb 2016

Process: 5896

Responsibility Allocation : Ensuring ORD's Are Taken To Goods Out And Invoices Are Retrieved 25 Feb 2016

Process: 5899

Proforma And Quote Chasing 25 Feb 2016

Process: 5901

Link Call Log Contacts To The CRM 02 Mar 2016

Process: 5913

Check For Humanmed Orders In Logistics Mailbox 03 Mar 2016

Process: 5943

Check Cardea And Multiquote 08 Mar 2016

Process: 5945

Responsibility Allocation : Sending Samples 08 Mar 2016

Process: 5946

Responsibility Allocation : Sending Sale Or Returns 08 Mar 2016

Process: 5947

Responsibility Allocation : Search For Distributors 08 Mar 2016

Process: 5948

Adding New Accounts To Opera 08 Mar 2016

Process: 5949

Filling Credit Card Slips 08 Mar 2016

Process: 6958

Responsibility Allocation : Shipped Order Queries 09 Mar 2016

Process: 7676

PDFing Of Invoices Viamed 17 Mar 2016

Process: 7693

Collect Repair Filing From Warehouse 22 Apr 2016

Process: 7752

SRS Folder 22 Nov 2016

Process: 7758

Check For GHX Orders 17 Jan 2017

Process: 7760

Send Service Offers 31 Jan 2017

Process: 7761

Send VST Delivery Notifications 01 Feb 2017

Process: 7795

Answering UK Web Questions 27 Apr 2017

Process: 7822

Review Oxylink Stock 26 Jul 2017

Process: 5859

Review Un-shipped Parcels 17 Feb 2016

Process: 6954

Back Orders Review - By Customer 09 Mar 2016

Process: 7748

		Check Repair Orders 10 Oct 2016 Process: 7749 Check Repair Quotes 10 Oct 2016
8.2.2 When determining the requirements for the products and services to be offered to customers, the organization shall ensure that: a) the requirements for the products and services are defined, including: 1) any applicable statutory and regulatory requirements; 2) those considered necessary by the organization; b) the organization can meet the claims for the products and services it offers. Determining the requirements for products and services	Top Level Document: VOP 03 Contract Review, Enquires, Office Processes Revision Document ID75949 **Date Revision 24 Nov 2021 Reviewed 24 Nov 2021 Top Level Document: VOP 17 Design Research and Development Revision Document ID25632 Date Revision 19 Mar 2018 Reviewed 19 Mar 2018 Audit 02 Contract Review and Sales Order Processing Revision Document ID69328 Date Revision 09 Sep 2021 Reviewed 09 Sep 2021 Audit 12 CE Files Revision Document ID63815 Date Revision 30 Jun 2021 Reviewed 30 Jun 2021 Audit 16 Sales and Marketing Revision Document ID69457 Date Revision 10 Sep 2021 Reviewed 10 Sep 2021	Process: 7703 Vandagraph Pay Pal Retrieve Funds 23 May 2016 Process: 7702 Responsibility Allocation : Vandagraph Pay Pay Issue Refund 23 May 2016 Process: 7396 Responsibility Allocation : VST Stock Meeting `Goods Out` Review 09 Mar 2016 Process: 7387 Responsibility Allocation : VST Stock Meeting Purchase Order Requirements 09 Mar 2016
8.2.3 Review of the requirements for products and services		Process: 7709 Delivered not Invoiced 28 Jun 2016 Process: 7702 Responsibility Allocation : Vandagraph Pay Pay Issue Refund 23 May 2016 Process: 7686 Thorough Checking Of Awaiting Action Tray 21 Apr 2016 Process: 7685 Repairs Ready For Invoice 18 Apr 2016 Process: 7683 Check Stock For Proforma 18 Apr 2016 Process: 7678

Check Catalog 360 Circle For Quotes And Orders 08 Apr 2016
Process: 7398
Responsibility Allocation : VST Stock Meeting UPS Shipping Fuel
Surcharge 09 Mar 2016
Process: 7396
Responsibility Allocation : VST Stock Meeting `Goods Out` Review
09 Mar 2016
Process: 7385
Responsibility Allocation : VST Stock Meeting Sales Forward Orders
Review 09 Mar 2016
Process: 6938
Responsibility Allocation : Customer Database Updates 09 Mar 2016
Process: 6956
Responsibility Allocation : Sales Order Issues 09 Mar 2016
Process: 7090
Responsibility Allocation : Office Procedures 09 Mar 2016
Process: 6898
GHX Web Pricing 09 Mar 2016
Process: 5871
Check Sale Or Returns 17 Feb 2016
Process: 5876
E.Commerce Cardea And Multiquote 17 Feb 2016
Process: 5872
Check Sale Or Returns Export 17 Feb 2016
Process: 2
Answering Telephones 16 Feb 2016
Process: 5
Responsibility Allocation : Processing Of Sales Orders 16 Feb 2016
Process: 6
Responsibility Allocation : Updating Contact Management System 16
Feb 2016
Process: 7
Responsibility Allocation : Checking Of Sales Orders 16 Feb 2016
Process: 8
Responsibility Allocation : Order And Status Liaison With Customers
16 Feb 2016
Process: 9
Distribution Of Faxes 16 Feb 2016
Process: 10

Distribution Of Emails 16 Feb 2016

Process: 11

Distribution Of Mail 16 Feb 2016

Process: 14

Fax Paper 16 Feb 2016

Process: 15

Filing and Archiving 16 Feb 2016

Process: 16

Responsibility Allocation : Photocopying 16 Feb 2016

Process: 21

Office Sales Projects 16 Feb 2016

Process: 36

Emailing Of Invoices 16 Feb 2016

Process: 5875

Check Paypal For Orders 17 Feb 2016

Process: 5879

Responsibility Allocation : Customer Returning Goods On Our UPS Account 18 Feb 2016

Process: 5882

Responsibility Allocation : Send Post To Humanmed 24 Feb 2016

Process: 5892

Checking EBay And Amazon For Orders And Messages 25 Feb 2016

Process: 5893

Answering Website Questions 25 Feb 2016

Process: 5894

Checking Of Active List 25 Feb 2016

Process: 5895

Responsibility Allocation : Completing Office Job List 25 Feb 2016

Process: 5896

Responsibility Allocation : Ensuring ORD's Are Taken To Goods Out And Invoices Are Retrieved 25 Feb 2016

Process: 5899

Proforma And Quote Chasing 25 Feb 2016

Process: 5901

Link Call Log Contacts To The CRM 02 Mar 2016

Process: 5913

Check For Humanmed Orders In Logistics Mailbox 03 Mar 2016

Process: 5943

Check Cardea And Multiquote 08 Mar 2016

Process: 5944

Responsibility Allocation : Chasing Lost Customers 08 Mar 2016

Process: 5945

Responsibility Allocation : Sending Samples 08 Mar 2016

Process: 5947

Responsibility Allocation : Search For Distributors 08 Mar 2016

Process: 5946

Responsibility Allocation : Sending Sale Or Returns 08 Mar 2016

Process: 5948

Adding New Accounts To Opera 08 Mar 2016

Process: 6958

Responsibility Allocation : Shipped Order Queries 09 Mar 2016

Process: 7676

PDFing Of Invoices Viamed 17 Mar 2016

Process: 7693

Collect Repair Filing From Warehouse 22 Apr 2016

Process: 7953

Vandagraph Delivery Notifications 26 May 2020

Process: 7699

Shred Sensitive Paperwork In JL Office 19 May 2016

Process: 7712

Review Inward Payments 01 Jul 2016

Process: 7735

Ensure SOR's Are Followed Up 01 Sep 2016

Process: 7752

SRS Folder 22 Nov 2016

Process: 7758

Check For GHX Orders 17 Jan 2017

Process: 7760

Send Service Offers 31 Jan 2017

Process: 7761

Send VST Delivery Notifications 01 Feb 2017

Process: 7783

PDF VST Invoices And Purchase Orders 10 Feb 2017

Process: 7792

Shipped Order Success Report 13 Mar 2017

Process: 7795

Answering UK Web Questions 27 Apr 2017

Process: 7822

		Review Oxylink Stock 26 Jul 2017 Process: 5859 Review Un-shipped Parcels 17 Feb 2016 Process: 6954 Back Orders Review - By Customer 09 Mar 2016 Process: 7749 Check Repair Quotes 10 Oct 2016 Process: 7748 Check Repair Orders 10 Oct 2016
8.2.3.1 The organization shall ensure that it has the ability to meet the requirements for products and services to be offered to customers. The organization shall conduct a review before committing to supply products and services to a customer, to include: a) requirements specified by the customer, including the requirements for delivery and postdelivery activities; b) requirements not stated by the customer, but necessary for the specified or intended use, when known; c) requirements specified by the organization; d) statutory and regulatory requirements applicable to the products and services; e) contract or order requirements differing from those previously expressed. The organization shall ensure that contract or order requirements differing from those previously defined are resolved. The customers requirements shall be confirmed by the organization before acceptance, when the	Top Level Document: VOP 03 Contract Review, Enquires, Office Processes Revision Document ID75949 **Date Revision 24 Nov 2021 Reviewed 24 Nov 2021 Audit 02 Contract Review and Sales Order Processing Revision Document ID69328 Date Revision 09 Sep 2021 Reviewed 09 Sep 2021	Process: 7831 Intrastats Debtors And Creditor Figures 18 Sep 2017 Process: 7796 Review Franking Label Errors 08 May 2017 Process: 7795 Answering UK Web Questions 27 Apr 2017 Process: 7749 Check Repair Quotes 10 Oct 2016 Process: 7748 Check Repair Orders 10 Oct 2016 Process: 7734 Responsibility Allocation : Humanmed Order Processing 25 Aug 2016 Process: 7712 Review Inward Payments 01 Jul 2016 Process: 7710 Responsibility Allocation : Proforma And Quote Processing 29 Jun 2016 Process: 7953 Vandagraph Delivery Notifications 26 May 2020 Process: 7691 Ship Sale Or Returns 21 Apr 2016 Process: 7684 Repairs Ready For Quote 18 Apr 2016 Process: 7674 Check Repairs Ready For Invoice List 10 Mar 2016 Process: 7390 Responsibility Allocation : VST Stock Meeting Returns Overview - Credits 09 Mar 2016 Process: 7387

customer does not provide a documented statement of their requirements. NOTE In some situations, such as internet sales, a formal review is impractical for each order. Instead, the review can cover relevant product information, such as catalogues.		Responsibility Allocation : VST Stock Meeting Purchase Order Requirements 09 Mar 2016
8.2.3.2 The organization shall retain documented information, as applicable: a) on the results of the review; b) on any new requirements for the products and services.	Top Level Document: VOP 01 Documentation and Records, Control, Creation, Storage, Retrieval, Revision Control and Online Records Revision Document ID75407 Date Revision 18 Nov 2021 Reviewed 18 Nov 2021 Audit 02 Contract Review and Sales Order Processing Revision Document ID69328 Date Revision 09 Sep 2021 Reviewed 09 Sep 2021 Audit 22 Post Market Surveillance Revision Document ID63052 Date Revision 22 Jun 2021 Reviewed 22 Jun 2021	Process: 7788 Petty Cash Reconciliation 02 Mar 2017 Process: 7674 Check Repairs Ready For Invoice List 10 Mar 2016
8.2.4 Changes to requirements for products and services The organization shall ensure that relevant documented information is amended, and that relevant persons are made aware of the changed requirements, when the requirements for products and services are changed.	Top Level Document: VOP 01 Documentation and Records, Control, Creation, Storage, Retrieval, Revision Control and Online Records Revision Document ID75407 Date Revision 18 Nov 2021 Reviewed 18 Nov 2021 Top Level Document: VOP 03 Contract Review, Enquires, Office Processes Revision Document ID75949 **Date Revision 24 Nov 2021 Reviewed 24 Nov 2021 Audit 02 Contract Review and Sales Order Processing Revision Document ID69328	Process: 7674 Check Repairs Ready For Invoice List 10 Mar 2016

	Date Revision 09 Sep 2021 Reviewed 09 Sep 2021 Audit 10 Documentation Control Revision Document ID63807 Date Revision 30 Jun 2021 Reviewed 30 Jun 2021	
8.3 Design and development of products and services	VM3COP02.01 Boundaries ISO 9001:2015 VST Revision Document ID69692 Date Revision 14 Sep 2021 Reviewed 14 Sep 2021	Process: 7810 Research Activities 06 Jun 2017
8.3.1 General The organization shall establish, implement and maintain a design and development process that is appropriate to ensure the subsequent provision of products and services.	Top Level Document: VOP 17 Design Research and Development Revision Document ID25632 Date Revision 19 Mar 2018 Reviewed 19 Mar 2018 Audit 03 Design Control Revision Document ID51631 Date Revision 13 Jan 2021 Reviewed 13 Jan 2021	
8.3.2 In determining the stages and controls for design and development, the organization shall consider: a) the nature, duration and complexity of the design and development activities; b) the required process stages, including applicable design and development reviews; c) the required design and development verification and validation activities; d) the responsibilities and authorities involved in the design and development process; e) the internal and external resource needs for the design and development of products and services; f) the need to control interfaces between	Top Level Document: VOP 17 Design Research and Development Revision Document ID25632 Date Revision 19 Mar 2018 Reviewed 19 Mar 2018 Audit 03 Design Control Revision Document ID51631 Date Revision 13 Jan 2021 Reviewed 13 Jan 2021 Audit 12 CE Files Revision Document ID63815 Date Revision 30 Jun 2021 Reviewed 30 Jun 2021 Audit 10 Documentation Control Revision Document ID63807 Date Revision 30 Jun 2021 Reviewed 30 Jun 2021	

persons involved in the design and development process; g) the need for involvement of customers and users in the design and development process; h) the requirements for subsequent provision of products and services; i) the level of control expected for the design and development process by customers and other relevant interested parties; j) the documented information needed to demonstrate that design and development requirements have been met. Design and development planning		
8.3.3 The organization shall determine the requirements essential for the specific types of products and services to be designed and developed. The organization shall consider: a) functional and performance requirements; b) information derived from previous similar design and development activities; c) statutory and regulatory requirements; d) standards or codes of practice that the organization has committed to implement; e) potential consequences of failure due to the nature of the products and services. Inputs shall be adequate for design and development purposes, complete and unambiguous. Conflicting design and development inputs shall be resolved. The organization shall retain documented information on design and development inputs. Design and development inputs	Top Level Document: VOP 17 Design Research and Development Revision Document ID25632 Date Revision 19 Mar 2018 Reviewed 19 Mar 2018 Audit 03 Design Control Revision Document ID51631 Date Revision 13 Jan 2021 Reviewed 13 Jan 2021 Audit 12 CE Files Revision Document ID63815 Date Revision 30 Jun 2021 Reviewed 30 Jun 2021 Audit 22 Post Market Surveillance Revision Document ID63052 Date Revision 22 Jun 2021 Reviewed 22 Jun 2021	Process: 7816 Repairs In Process Review 06 Jun 2017 Process: 7814 Responsibility Allocation : Viamed Repairs 06 Jun 2017 Process: 7744 FDA Device Establishment Registration And Listing 28 Sep 2016 Process: 7705 Checking For Uploaded Files 08 Jun 2016

8.3.4 The organization shall apply controls to the design and development process to ensure that: a) the results to be achieved are defined; b) reviews are conducted to evaluate the ability of the results of design and development to meet requirements; c) verification activities are conducted to ensure that the design and development outputs meet the input requirements; d) validation activities are conducted to ensure that the resulting products and services meet the requirements for the specified application or intended use; e) any necessary actions are taken on problems determined during the reviews, or verification and validation activities; f) documented information of these activities is retained. NOTE Design and development reviews, verification and validation have distinct purposes. They can be conducted separately or in any combination, as is suitable for the products and services of the organization. Design and development controls	Top Level Document: VOP 17 Design Research and Development Revision Document ID25632 Date Revision 19 Mar 2018 Reviewed 19 Mar 2018 Audit 03 Design Control Revision Document ID51631 Date Revision 13 Jan 2021 Reviewed 13 Jan 2021 Audit 10 Documentation Control Revision Document ID63807 Date Revision 30 Jun 2021 Reviewed 30 Jun 2021 Audit 22 Post Market Surveillance Revision Document ID63052 Date Revision 22 Jun 2021 Reviewed 22 Jun 2021	
8.3.5 The organization shall ensure that design and development outputs: a) meet the input requirements; b) are adequate for the subsequent processes for the provision of products and services; c) include or reference monitoring and	Top Level Document: VOP 17 Design Research and Development Revision Document ID25632 Date Revision 19 Mar 2018 Reviewed 19 Mar 2018 Audit 03 Design Control Revision Document ID51631 Date Revision 13 Jan 2021 Reviewed 13	Process: 7705 Checking For Uploaded Files 08 Jun 2016

measuring requirements, as appropriate, and acceptance criteria; d) specify the characteristics of the products and services that are essential for their intended purpose and their safe and proper provision. The organization shall retain documented information on design and development outputs. Design and development outputs	Jan 2021 Audit 10 Documentation Control Revision Document ID63807 Date Revision 30 Jun 2021 Reviewed 30 Jun 2021	
8.3.6 The organization shall identify, review and control changes made during, or subsequent to, the design and development of products and services, to the extent necessary to ensure that there is no adverse impact on conformity to requirements. The organization shall retain documented information on: a) design and development changes; b) the results of reviews; c) the authorization of the changes; d) the actions taken to prevent adverse impacts. Design and development changes	Top Level Document: VOP 17 Design Research and Development Revision Document ID25632 Date Revision 19 Mar 2018 Reviewed 19 Mar 2018 Audit 03 Design Control Revision Document ID51631 Date Revision 13 Jan 2021 Reviewed 13 Jan 2021 Audit 20 Process verification to Managment Revision Document ID73324 Date Revision 26 Oct 2021 Reviewed 26 Oct 2021 Audit 22 Post Market Surveillance Revision Document ID63052 Date Revision 22 Jun 2021 Reviewed 22 Jun 2021	Process: 7830 Review Q.A. Failures Report 18 Sep 2017 Process: 7705 Checking For Uploaded Files 08 Jun 2016
8.4 Control of externally provided processes, products and services	VM3COP02.01 Boundaries ISO 9001:2015 VST Revision Document ID69692 Date Revision 14 Sep 2021 Reviewed 14 Sep 2021	Process: 7707 Send Purchase Orders To Suppliers 13 Jun 2016 Process: 7682 Check Stock Requirements Supplier Bluepoint 18 Apr 2016 Process: 7681 Check Stock Requirements Supplier Posey 18 Apr 2016 Process: 7680 Check Stock Requirements Supplier Envitec 18 Apr 2016 Process: 7679 Check Stock Requirements Supplier Teledyne 18 Apr 2016 Process: 7675

		Responsibility Allocation : Ordering Demo Stock For Humanmed Reps 11 Mar 2016 Process: 7395 Responsibility Allocation : VST Stock Meeting `Goods In` Review 09 Mar 2016
8.4.1 The organization shall ensure that externally provided processes, products and services conform to requirements. The organization shall determine the controls to be applied to externally provided processes, products and services when: a) products and services from external providers are intended for incorporation into the organization's own products and services; b) products and services are provided directly to the customer(s) by external providers on behalf of the organization; c) a process, or part of a process, is provided by an external provider as a result of a decision by the organization. The organization shall determine and apply criteria for the evaluation, selection, monitoring of performance, and re-evaluation of external providers, based on their ability to provide processes or products and services in accordance with requirements. The organization shall retain documented information of these activities and any necessary actions arising from the evaluations. General	Top Level Document: VOP 05 Supplier Control, Supplier Review, Purchase Orders, Supplier Returns and Rejection Revision Document ID75847 **Date Revision 23 Nov 2021 Reviewed 23 Nov 2021 Audit 05 Purchasing suppliers Revision Document ID69314 Date Revision 09 Sep 2021 Reviewed 09 Sep 2021 Audit 07 Handling and Storage Revision Document ID58347 Date Revision 23 Apr 2021 Reviewed 23 Apr 2021	Process: 7826 Goods In Processes 06 Sep 2017 Process: 7799 Opera Purchase Ledger Close 11 May 2017 Process: 7755 Fast Hosts Invoice 08 Dec 2016 Process: 7701 AWS Amazon Web Services 23 May 2016 Process: 7700 Domain Name Management 19 May 2016 Process: 7387 Responsibility Allocation : VST Stock Meeting Purchase Order Requirements 09 Mar 2016 Process: 7707 Send Purchase Orders To Suppliers 13 Jun 2016

8.4.2 The organization shall ensure that externally provided processes, products and services do not adversely affect the organization's ability to consistently deliver conforming products and services to its customers. The organization shall: a) ensure that externally provided processes remain within the control of its quality management system; b) define both the controls that it intends to apply to an external provider and those it intends to apply to the resulting output; c) take into consideration: 1) the potential impact of the externally provided processes, products and services on the organization's ability to consistently meet customer and applicable statutory and regulatory requirements; 2) the effectiveness of the controls applied by the external provider; d) determine the verification, or other activities, necessary to ensure that the externally provided processes, products and services meet requirements. Type and extent of control	Top Level Document: VOP 05 Supplier Control, Supplier Review, Purchase Orders, Supplier Returns and Rejection Revision Document ID75847 **Date Revision 23 Nov 2021 Reviewed 23 Nov 2021 Audit 05 Purchasing suppliers Revision Document ID69314 Date Revision 09 Sep 2021 Reviewed 09 Sep 2021	Process: 7826 Goods In Processes 06 Sep 2017 Process: 7751 VST Purchase Order Log 02 Nov 2016
8.4.3 The organization shall ensure the adequacy of requirements prior to their communication to the external provider. The organization shall communicate to external providers its requirements for:	Top Level Document: VOP 05 Supplier Control, Supplier Review, Purchase Orders, Supplier Returns and Rejection Revision Document ID75847 **Date Revision 23 Nov 2021 Reviewed 23 Nov 2021 Audit 05 Purchasing suppliers	Process: 7826 Goods In Processes 06 Sep 2017 Process: 7823 Safety Tester Data 02 Aug 2017 Process: 7787 Check Returns All Supplier 15 Feb 2017 Process: 7786

a) the processes, products and services to be provided; b) the approval of: 1) products and services; 2) methods, processes and equipment; 3) the release of products and services; c) competence, including any required qualification of persons; d) the external providers interactions with the organization; e) control and monitoring of the external providers performance to be applied by the organization; f) verification or validation activities that the organization, or its customer, intends to perform at the external providers premises. Information for external providers	Revision Document ID69314 Date Revision 09 Sep 2021 Reviewed 09 Sep 2021	Check Returns Supplier Maxtec 15 Feb 2017 Process: 7785 Check Returns Supplier Teledyne 15 Feb 2017 Process: 7784 Check Returns Supplier Envitec 15 Feb 2017 Process: 7387 Responsibility Allocation : VST Stock Meeting Purchase Order Requirements 09 Mar 2016
8.5 Production and service provision		Process: 7738 Production Statistics 03 Sep 2016
8.5.1 The organization shall implement production and service provision under controlled conditions. Controlled conditions shall include, as applicable: a) the availability of documented information that defines: 1) the characteristics of the products to be produced, the services to be provided, or the activities to be performed; 2) the results to be achieved; b) the availability and use of suitable monitoring and measuring resources; c) the implementation of monitoring and measurement activities at appropriate	Top Level Document: VOP 08 Production, Reworks, New Production Revision Document ID31072 Date Revision 30 Sep 2019 Reviewed 30 Sep 2019 Top Level Document: VOP 07 Stock Control, Handling, Control of Labelling, Storage, Movement Revision Document ID75973 **Date Revision 24 Nov 2021 Reviewed 24 Nov 2021 Top Level Document: VOP 06 Measurement Control Viamed VST, Calibration, QA Stock Revision Document ID53615 Date Revision 11 Feb 2021 Reviewed 11 Feb 2021	Process: 7737 Production In Production List 03 Sep 2016 Process: 7736 Production Start Job List 03 Sep 2016 Process: 7682 Check Stock Requirements Supplier Bluepoint 18 Apr 2016 Process: 7681 Check Stock Requirements Supplier Posey 18 Apr 2016 Process: 7680 Check Stock Requirements Supplier Envitec 18 Apr 2016 Process: 7679 Check Stock Requirements Supplier Teledyne 18 Apr 2016 Process: 7675 Responsibility Allocation : Ordering Demo Stock For Humanmed Reps 11 Mar 2016 Process: 7401 Responsibility Allocation : VST Calibration 09 Mar 2016

stages to verify that criteria for control of processes or outputs, and acceptance criteria for products and services, have been met;
d) the use of suitable infrastructure and environment for the operation of processes;
e) the appointment of competent persons, including any required qualification;
f) the validation, and periodic revalidation, of the ability to achieve planned results of the processes for production and service provision, where the resulting output cannot be verified by subsequent monitoring or measurement;
g) the implementation of actions to prevent human error;
h) the implementation of release, delivery and post-delivery activities **Control of production and service provision**

Top Level Document: VOP 22 Picking and Packing Dispatch and Goods Out
Revision Document ID31048
Date Revision 30 Sep 2019 Reviewed 30 Sep 2019
Top Level Document: VOP 27 Software Validation
Revision Document ID31064
Date Revision 30 Sep 2019 Reviewed 30 Sep 2019
Top Level Document: VOP 02 Personnel and Responsibility , Staff and Staffing Issues, Training, Roles and Tasks
Revision Document ID73529
Date Revision 29 Oct 2021 Reviewed 29 Oct 2021
Audit 03 Design Control
Revision Document ID51631
Date Revision 13 Jan 2021 Reviewed 13 Jan 2021
Audit 07 Handling and Storage
Revision Document ID58347
Date Revision 23 Apr 2021 Reviewed 23 Apr 2021
Audit 08 Training, Competence and Human Resources
Revision Document ID70147
Date Revision 20 Sep 2021 Reviewed 20 Sep 2021
Audit 24 Service Logs
Revision Document ID68263
Date Revision 26 Aug 2021 Reviewed 26 Aug 2021
Audit 06 Calibration
Revision Document ID63048
Date Revision 22 Jun 2021 Reviewed 22 Jun 2021
VM3COP20.37 Generating a New Service Visit

Process: 7395
Responsibility Allocation : VST Stock Meeting `Goods In` Review
09 Mar 2016
Process: 7048
Control of monitoring and measuring devices 09 Mar 2016

	Revision Document ID17116 Date Revision 28 Jun 2016 Reviewed 28 Jun 2016 Audit 07 Handling and Storage Revision Document ID58347 Date Revision 23 Apr 2021 Reviewed 23 Apr 2021 Audit 15 Production Revision Document ID59614 Date Revision 11 May 2021 Reviewed 11 May 2021 Audit 09 Goods Inward and Product Identity Revision Document ID55437 Date Revision 12 Mar 2021 Reviewed 12 Mar 2021 Audit 01 Picking packing Revision Document ID51629 Date Revision 13 Jan 2021 Reviewed 13 Jan 2021	
8.5.2 The organization shall use suitable means to identify outputs when it is necessary to ensure the conformity of products and services. The organization shall identify the status of outputs with respect to monitoring and measurement requirements throughout production and service provision. The organization shall control the unique identification of the outputs when traceability is a requirement, and shall retain the documented information necessary to enable traceability. Identification and traceability	Top Level Document: VOP 09 Repairs and Servicing Revision Document ID75927 **Date Revision 24 Nov 2021 Reviewed 24 Nov 2021 Top Level Document: VOP 20 Goods in Purchases, Returns, Repairs, Inspection / Rejection Revision Document ID75943 **Date Revision 24 Nov 2021 Reviewed 24 Nov 2021 Audit 07 Handling and Storage Revision Document ID58347 Date Revision 23 Apr 2021 Reviewed 23 Apr 2021 Audit 09 Goods Inward and Product Identity Revision Document ID55437	Process: 7830 Review Q.A. Failures Report 18 Sep 2017 Process: 7737 Production In Production List 03 Sep 2016 Process: 7682 Check Stock Requirements Supplier Bluepoint 18 Apr 2016 Process: 7681 Check Stock Requirements Supplier Posey 18 Apr 2016 Process: 7680 Check Stock Requirements Supplier Envitec 18 Apr 2016 Process: 7679 Check Stock Requirements Supplier Teledyne 18 Apr 2016 Process: 7675 Responsibility Allocation : Ordering Demo Stock For Humanmed Reps 11 Mar 2016 Process: 7395 Responsibility Allocation : VST Stock Meeting `Goods In` Review 09 Mar 2016

	Date Revision 12 Mar 2021 Reviewed 12 Mar 2021	
8.5.3 The organization shall exercise care with property belonging to customers or external providers while it is under the organization's control or being used by the organization. The organization shall identify, verify, protect and safeguard customers' or external providers' property provided for use or incorporation into the products and services. When the property of a customer or external provider is lost, damaged or otherwise found to be unsuitable for use, the organization shall report this to the customer or external provider and retain documented information on what has occurred. NOTE A customer's or external provider's property can include materials, components, tools and equipment, premises, intellectual property and personal data. Property belonging to customers or external providers	Top Level Document: VOP 09 Repairs and Servicing Revision Document ID75927 **Date Revision 24 Nov 2021 Reviewed 24 Nov 2021 Top Level Document: VOP 20 Goods in Purchases, Returns, Repairs, Inspection / Rejection Revision Document ID75943 **Date Revision 24 Nov 2021 Reviewed 24 Nov 2021 Audit 07 Handling and Storage Revision Document ID58347 Date Revision 23 Apr 2021 Reviewed 23 Apr 2021 Audit 11 Repairs, Servicing and Returns Revision Document ID64142 Date Revision 02 Jul 2021 Reviewed 02 Jul 2021 Audit 09 Goods Inward and Product Identity Revision Document ID55437 Date Revision 12 Mar 2021 Reviewed 12 Mar 2021	Process: 7823 Saftey Tester Data 02 Aug 2017 Process: 7814 Responsibility Allocation : Viamed Repairs 06 Jun 2017 Process: 7813 Responsibility Allocation : VST Repairs 06 Jun 2017 Process: 7812 Responsibility Allocation : Vandagraph Repairs 06 Jun 2017 Process: 7735 Ensure SOR's Are Followed Up 01 Sep 2016
8.5.4 The organization shall preserve the outputs during production and service provision, to the extent necessary to ensure conformity to requirements. NOTE Preservation can include identification, handling, contamination control, packaging, storage,	Top Level Document: VOP 07 Stock Control, Handling, Control of Labelling, Storage, Movement Revision Document ID75973 **Date Revision 24 Nov 2021 Reviewed 24 Nov 2021 Top Level Document: VM3COP27.51 Incoming / Goods in Contamination Control Revision Document ID74855	Process: 7830 Review Q.A. Failures Report 18 Sep 2017

transmission or transportation, and protection. Preservation	Date Revision 12 Nov 2021 Reviewed 12 Nov 2021 Audit 07 Handling and Storage Revision Document ID58347 Date Revision 23 Apr 2021 Reviewed 23 Apr 2021 Audit 09 Goods Inward and Product Identity Revision Document ID55437 Date Revision 12 Mar 2021 Reviewed 12 Mar 2021	
8.5.5 The organization shall meet requirements for post-delivery activities associated with the products and services. In determining the extent of post-delivery activities that are required, the organization shall consider: a) statutory and regulatory requirements; b) the potential undesired consequences associated with its products and services; c) the nature, use and intended lifetime of its products and services; d) customer requirements; e) customer feedback. NOTE Post-delivery activities can include actions under warranty provisions, contractual obligations such as maintenance services, and supplementary services such as recycling or final disposal. Post-delivery activities	Top Level Document: VOP 13 Process Monitoring, System Reviews, Audits, Management Reviews and Analysis Data Revision Document ID75461 Date Revision 18 Nov 2021 Reviewed 18 Nov 2021 Audit 20 Process verification to Managment Revision Document ID73324 Date Revision 26 Oct 2021 Reviewed 26 Oct 2021 Audit 14 Complaints and Corrective Actions Revision Document ID76091 **Date Revision 25 Nov 2021 Reviewed 25 Nov 2021 Audit 22 Post Market Surveillance Revision Document ID63052 Date Revision 22 Jun 2021 Reviewed 22 Jun 2021	Process: 7826 Goods In Processes 06 Sep 2017 Process: 7821 Controlled Waste Description And Transfer 15 Jun 2017 Process: 7820 North Yorkshire Council Waste Tranfer 15 Jun 2017 Process: 7735 Ensure SOR's Are Followed Up 01 Sep 2016 Process: 7427 Responsibility Allocation : VST Customer Complaints 09 Mar 2016 Process: 7391 Responsibility Allocation : VST Stock Meeting Customer Complaints Review **Mandatory** 09 Mar 2016 Process: 7389 Responsibility Allocation : VST Stock Meeting Returns Overview - From Customers 09 Mar 2016 Process: 7843 Review VST Product Feedback Negative 23 Sep 2017 Process: 7842 Review VIAMED Product Feedback Negative 23 Sep 2017 Process: 7841 Review VST Feedback - Customer Complaints 23 Sep 2017 Process: 7840 Review VST Feedback - Customer Feedback Negative 23 Sep 2017 Process: 7839 Review VIAMED Feedback - Customer Complaints 23 Sep 2017 Process: 7838

		Review VIAMED Feedback - Customer Feedback Negative 23 Sep 2017
8.5.6 The organization shall review and control changes for production or service provision, to the extent necessary to ensure continuing conformity with requirements. The organization shall retain documented information describing the results of the review of changes, the person(s) authorizing the change, and any necessary actions arising from the review. Control of changes	Audit 12 CE Files Revision Document ID63815 Date Revision 30 Jun 2021 Reviewed 30 Jun 2021	
8.6 The organization shall implement planned arrangements, at appropriate stages, to verify that the product and service requirements have been met. The release of products and services to the customer shall not proceed until the planned arrangements have been satisfactorily completed, unless otherwise approved by a relevant authority and, as applicable, by the customer. The organization shall retain documented information on the release of products and services. The documented information shall include: a) evidence of conformity with the acceptance criteria; b) traceability to the person(s) authorizing the release Release of products and services	Top Level Document: VOP 22 Picking and Packing Dispatch and Goods Out Revision Document ID31048 Date Revision 30 Sep 2019 Reviewed 30 Sep 2019 Audit 07 Handling and Storage Revision Document ID58347 Date Revision 23 Apr 2021 Reviewed 23 Apr 2021 Audit 09 Goods Inward and Product Identity Revision Document ID55437 Date Revision 12 Mar 2021 Reviewed 12 Mar 2021	Process: 7830 Review Q.A. Failures Report 18 Sep 2017
8.7 Control of nonconforming outputs		Process: 7671 Humanmed Non Conformances 09 Mar 2016

8.7.1 The organization shall ensure that outputs that do not conform to their requirements are identified and controlled to prevent their unintended use or delivery. The organization shall take appropriate action based on the nature of the nonconformity and its effect on the conformity of products and services. This shall also apply to nonconforming products and services detected after delivery of products, during or after the provision of services. The organization shall deal with nonconforming outputs in one or more of the following ways: - correction; - segregation, containment, return or suspension of provision of products and services; - informing the customer; - obtaining authorization for acceptance under concession. Conformity to the requirements shall be verified when nonconforming outputs are corrected.	Top Level Document: VOP 07 Stock Control, Handling, Control of Labelling, Storage, Movement Revision Document ID75973 **Date Revision 24 Nov 2021 Reviewed 24 Nov 2021 Top Level Document: VOP 06 Measurement Control Viamed VST, Calibration, QA Stock Revision Document ID53615 Date Revision 11 Feb 2021 Reviewed 11 Feb 2021 Audit 05 Purchasing suppliers Revision Document ID69314 Date Revision 09 Sep 2021 Reviewed 09 Sep 2021 Audit 07 Handling and Storage Revision Document ID58347 Date Revision 23 Apr 2021 Reviewed 23 Apr 2021 Audit 09 Goods Inward and Product Identity Revision Document ID55437 Date Revision 12 Mar 2021 Reviewed 12 Mar 2021	Process: 7830 Review Q.A. Failures Report 18 Sep 2017 Process: 7826 Goods In Processes 06 Sep 2017 Process: 7752 SRS Folder 22 Nov 2016 Process: 7749 Check Repair Quotes 10 Oct 2016 Process: 7690 Ship Repairs 21 Apr 2016 Process: 7685 Repairs Ready For Invoice 18 Apr 2016 Process: 7684 Repairs Ready For Quote 18 Apr 2016 Process: 7674 Check Repairs Ready For Invoice List 10 Mar 2016 Process: 7671 Humanmed Non Conformances 09 Mar 2016 Process: 7399 Responsibility Allocation : VST Stock Meeting Non Conforming Stock Transfers. (QC19) 09 Mar 2016 Process: 7394 Responsibility Allocation : VST Stock Meeting Repairs Review - General 09 Mar 2016 Process: 7390 Responsibility Allocation : VST Stock Meeting Returns Overview - Credits 09 Mar 2016 Process: 7388 Responsibility Allocation : VST Stock Meeting Returns Overview 09 Mar 2016
8.7.2 The organization shall retain documented information that: - describes the nonconformity; - describes the actions taken; - describes any concessions obtained; - identifies the authority deciding the action in respect of the nonconformity.	Audit 20 Process verification to Managment Revision Document ID73324 Date Revision 26 Oct 2021 Reviewed 26 Oct 2021 Audit 12 CE Files Revision Document ID63815 Date Revision 30 Jun 2021 Reviewed 30 Jun 2021	Process: 7830 Review Q.A. Failures Report 18 Sep 2017 Process: 7690 Ship Repairs 21 Apr 2016 Process: 7671 Humanmed Non Conformances 09 Mar 2016 Process: 7394 Responsibility Allocation : VST Stock Meeting Repairs Review - General 09 Mar 2016

9 Performance evaluation

9 Performance evaluation		Process: 7433 Responsibility Allocation : VST Board Directors Meeting 09 Mar 2016
9.1 Monitoring, measurement, analysis and evaluation		
9.1.1 The organization shall determine: a) what needs to be monitored and measured; b) the methods for monitoring, measurement, analysis and evaluation needed to ensure valid results; c) when the monitoring and measuring shall be performed; d) when the results from monitoring and measurement shall be analysed and evaluated. The organization shall evaluate the performance and the effectiveness of the quality management system. The organization shall retain appropriate documented information as evidence of the results. General	Top Level Document: VOP 10 Non Conformance, Corrective and Preventive Actions Revision Document ID46915 Date Revision 02 Nov 2020 Reviewed 02 Nov 2020 Top Level Document: VOP 13 Process Monitoring, System Reviews, Audits, Management Reviews and Analysis Data Revision Document ID75461 Date Revision 18 Nov 2021 Reviewed 18 Nov 2021 Audit 10 Documentation Control Revision Document ID63807 Date Revision 30 Jun 2021 Reviewed 30 Jun 2021 Audit 07 Handling and Storage Revision Document ID58347 Date Revision 23 Apr 2021 Reviewed 23 Apr 2021	Process: 7693 Collect Repair Filing From Warehouse 22 Apr 2016 Process: 7692 Responsibility Allocation : Take Complete Repair Paperwork To Office 22 Apr 2016 Process: 7394 Responsibility Allocation : VST Stock Meeting Repairs Review - General 09 Mar 2016
9.1.2 The organization shall monitor customers' perceptions of the degree to which their needs and expectations have been fulfilled. The organization shall determine the methods for obtaining,	Top Level Document: VOP 13 Process Monitoring, System Reviews, Audits, Management Reviews and Analysis Data Revision Document ID75461 Date Revision 18 Nov 2021 Reviewed 18 Nov 2021	Process: 7825 Responsibility Allocation : Order Picking 06 Sep 2017 Process: 7822 Review Oxylink Stock 26 Jul 2017 Process: 7797 Check Order Are Being Picked In Priority Order 10 May 2017 Process: 7693

monitoring and reviewing this information. NOTE Examples of monitoring customer perceptions can include customer surveys, customer feedback on delivered products and services, meetings with customers, market-share analysis, compliments, warranty claims and dealer reports. Customer satisfaction	Audit 14 Complaints and Corrective Actions Revision Document ID76091 **Date Revision 25 Nov 2021 Reviewed 25 Nov 2021 Audit 22 Post Market Surveillance Revision Document ID63052 Date Revision 22 Jun 2021 Reviewed 22 Jun 2021	Collect Repair Filing From Warehouse 22 Apr 2016 Process: 7692 Responsibility Allocation : Take Complete Repair Paperwork To Office 22 Apr 2016 Process: 7673 Check Expiry Dated Stock 09 Mar 2016 Process: 7664 Responsibility Allocation : Marketing Job Logger 09 Mar 2016 Process: 7427 Responsibility Allocation : VST Customer Complaints 09 Mar 2016 Process: 7394 Responsibility Allocation : VST Stock Meeting Repairs Review - General 09 Mar 2016 Process: 7391 Responsibility Allocation : VST Stock Meeting Customer Complaints Review **Mandatory** 09 Mar 2016 Process: 7389 Responsibility Allocation : VST Stock Meeting Returns Overview - From Customers 09 Mar 2016 Process: 7843 Review VST Product Feedback Negative 23 Sep 2017 Process: 7842 Review VIAMED Product Feedback Negative 23 Sep 2017 Process: 7841 Review VST Feedback - Customer Complaints 23 Sep 2017 Process: 7840 Review VST Feedback - Customer Feedback Negative 23 Sep 2017 Process: 7839 Review VIAMED Feedback - Customer Complaints 23 Sep 2017 Process: 7838 Review VIAMED Feedback - Customer Feedback Negative 23 Sep 2017
9.1.3 The organization shall analyse and evaluate appropriate data and information arising from monitoring and measurement. The results of analysis shall be used to evaluate:	Top Level Document: VOP 13 Process Monitoring, System Reviews, Audits, Management Reviews and Analysis Data Revision Document ID75461 Date Revision 18 Nov 2021 Reviewed 18 Nov 2021	Process: 7830 Review Q.A. Failures Report 18 Sep 2017 Process: 7822 Review Oxylink Stock 26 Jul 2017 Process: 7394 Responsibility Allocation : VST Stock Meeting Repairs Review - General 09 Mar 2016

- a) conformity of products and services;
- b) the degree of customer satisfaction;
- c) the performance and effectiveness of the quality management system;
- d) if planning has been implemented effectively;
- e) the effectiveness of actions taken to address risks and opportunities;
- f) the performance of external providers;
- g) the need for improvements to the quality management system.

NOTE Methods to analyse data can include statistical techniques. **Analysis and evaluation**

Top Level Document: VOP 05 Supplier Control, Supplier Review, Purchase Orders, Supplier Returns and Rejection
Revision Document ID75847

****Date Revision 23 Nov 2021 Reviewed 23 Nov 2021**

Audit 22 Post Market Surveillance
Revision Document ID63052

Date Revision 22 Jun 2021 Reviewed 22 Jun 2021

Process: 27

Management Reviews And Quality Audits 16 Feb 2016

Process: 7834

Financial Review 20 Sep 2017

Process: 26

Company Resources 16 Feb 2016

Process: 7713

Review Roles And Responsibilitys 17 Aug 2016

Process: 7837

Review External Parties Influencing The QMS VST / Viamed 23 Sep 2017

Process: 7840

Review VST Feedback - Customer Feedback Negative 23 Sep 2017

Process: 7841

Review VST Feedback - Customer Complaints 23 Sep 2017

Process: 7843

Review VST Product Feedback Negative 23 Sep 2017

Process: 7846

ISO System Management Review Viamed 26 Sep 2017

Process: 7848

Review ISO Scopes 27 Sep 2017

Process: 7849

Review Product Failures New Codes 28 Sep 2017

Process: 7871

Review Exclusion From Viamed 13485:2016 And VST 9001:2015 15 Oct 2017

Process: 7876

Maintain Update Of ISO Route Maps 21 Oct 2017

Process: 7878

Review Possible Upcoming Regulation Changes 22 Oct 2017

Process: 28

Supplier Review 16 Feb 2016

Process: 5889

Responsibility Allocation : Audit And Task - Audit 24 Feb 2016

Process: 7071

Post Market Surveillance 09 Mar 2016

Process: 7199

Non Conformities Review Viamed 09 Mar 2016

Process: 7743

		Customer Complaints Paper File 26 Sep 2016 Process: 7827 Review The Quality Policy VST 16 Sep 2017 Process: 7833 Importance Of Effective Quality Management 20 Sep 2017 Process: 6829 Supplier Review - Outstanding orders 09 Mar 2016 Process: 6832 Supplier Review Future orders 09 Mar 2016 Process: 7091 Calibration Index 09 Mar 2016 Process: 5881 Training Records Review 18 Feb 2016 Process: 7847 Health And Safety Review 26 Sep 2017 Process: 7793 Team Review Meeting 16 Mar 2017
9.2 Internal audi		Process: 7781 Audit 23 Analysis Of Data VST 08 Feb 2017 Process: 7780 Audit 22 Post Market Surveillance VST 08 Feb 2017 Process: 7779 Audit 21 Audit Of Audit VST 08 Feb 2017 Process: 7778 Audit 20 Process Verification To Managment VST 08 Feb 2017 Process: 7777 Audit 19 Health And Saftey VST 08 Feb 2017 Process: 7776 Audit 17 Internal Audits VST 08 Feb 2017 Process: 7775 Audit 15 Production VST 08 Feb 2017 Process: 7774 Audit 14 Complaints And Corrective Actions VST 08 Feb 2017 Process: 7773 Audit 12 CE Files VST 08 Feb 2017 Process: 7772 Audit 11 Repairs And Service VST 08 Feb 2017 Process: 7771 Audit 10b Process Verification VST 08 Feb 2017

Process: 7770

Audit 10 Documentation Control VST 08 Feb 2017

Process: 7769

Audit 09 Goods Inward And Product Identity VST 08 Feb 2017

Process: 7768

Audit 08 Training VST 08 Feb 2017

Process: 7767

Audit 07 Handling And Storage VST 08 Feb 2017

Process: 7766

Audit 06 Calibration VST 08 Feb 2017

Process: 7765

Audit 05 Purchasing Suppliers VST 08 Feb 2017

Process: 7764

Audit 03 Design Control VST 08 Feb 2017

Process: 7763

Audit 02 Contract Review VST 08 Feb 2017

Process: 7762

Audit 01 Picking Packing VST 08 Feb 2017

Process: 7733

Audit 23 Analysis Of Data Viamed 24 Aug 2016

Process: 7732

Audit 22 Post Market Surveillance Viamed 24 Aug 2016

Process: 7731

Audit 21 Audit Of Audit Viamed 24 Aug 2016

Process: 7730

Audit 20 Process Verification To Managment Viamed 24 Aug 2016

Process: 7729

Audit 19 Health And Saftey Viamed 24 Aug 2016

Process: 7728

Audit 17 Internal Audits Viamed 24 Aug 2016

Process: 7727

Audit 15 Production Viamed 24 Aug 2016

Process: 7726

Audit 14 Complaints And Corrective Actions Viamed 24 Aug 2016

Process: 7725

Audit 12 CE Files Viamed 24 Aug 2016

Process: 7724

Audit 11 Repairs And Service Viamed 24 Aug 2016

Process: 7723

		Audit 10b Process Verification Viamed 24 Aug 2016 Process: 7722 Audit 10 Documentation Control Viamed 24 Aug 2016 Process: 7721 Audit 09 Goods Inward And Product Identity Viamed 24 Aug 2016 Process: 7720 Audit 08 Training Viamed 24 Aug 2016 Process: 7719 Audit 07 Handling And Storage Viamed 24 Aug 2016 Process: 7718 Audit 06 Calibration Viamed 24 Aug 2016 Process: 7717 Audit 05 Purchasing Suppliers Viamed 24 Aug 2016 Process: 7716 Audit 03 Design Control Viamed 24 Aug 2016 Process: 7715 Audit 02 Contract Review Viamed 24 Aug 2016 Process: 7714 Audit 01 Picking Packing Viamed 24 Aug 2016
9.2.1 The organization shall conduct internal audits at planned intervals to provide information on whether the quality management system: a) conforms to: 1) the organization's own requirements for its quality management system; 2) the requirements of this International Standard; b) is effectively implemented and maintained.	Top Level Document: VOP 13 Process Monitoring, System Reviews, Audits, Management Reviews and Analysis Data Revision Document ID75461 Date Revision 18 Nov 2021 Reviewed 18 Nov 2021 Audit 20 Process verification to Managment Revision Document ID73324 Date Revision 26 Oct 2021 Reviewed 26 Oct 2021	Process: 7744 FDA Device Establishment Registration And Listing 28 Sep 2016 Process: 7668 Responsibility Allocation : Upgrading Intrastats ISO Quality system 09 Mar 2016
9.2.2 The organization shall: a) plan, establish, implement and maintain an audit programme(s) including the frequency, methods, responsibilities, planning requirements and reporting, which shall take into	Top Level Document: VOP 13 Process Monitoring, System Reviews, Audits, Management Reviews and Analysis Data Revision Document ID75461 Date Revision 18 Nov 2021 Reviewed 18 Nov 2021	

consideration the importance of the processes concerned, changes affecting the organization, and the results of previous audits; b) define the audit criteria and scope for each audit; c) select auditors and conduct audits to ensure objectivity and the impartiality of the audit process; d) ensure that the results of the audits are reported to relevant management; e) take appropriate correction and corrective actions without undue delay; f) retain documented information as evidence of the implementation of the audit programme and the audit results. NOTE See ISO 19011 for guidance.	Audit 10 Documentation Control Revision Document ID63807 Date Revision 30 Jun 2021 Reviewed 30 Jun 2021 Audit 18 Management Review Revision Document ID73320 **Date Revision 26 Oct 2021 Reviewed 26 Oct 2021	
9.3 Management review		
9.3.1 Top management shall review the organization's quality management system, at planned intervals, to ensure its continuing suitability, adequacy, effectiveness and alignment with the strategic direction of the organization. General	Top Level Document: VOP 13 Process Monitoring, System Reviews, Audits, Management Reviews and Analysis Data Revision Document ID75461 Date Revision 18 Nov 2021 Reviewed 18 Nov 2021 Audit 18 Management Review Revision Document ID73320 **Date Revision 26 Oct 2021 Reviewed 26 Oct 2021	
9.3.2 9.3.2 Management review inputs The management review shall be planned and carried out taking into consideration: a) the status of actions from previous management reviews;	Top Level Document: VOP 13 Process Monitoring, System Reviews, Audits, Management Reviews and Analysis Data Revision Document ID75461 Date Revision 18 Nov 2021 Reviewed 18	Process: 7831 Intrastats Debtors And Creditor Figures 18 Sep 2017 Process: 7830 Review Q.A. Failures Report 18 Sep 2017 Process: 7825 Responsibility Allocation : Order Picking 06 Sep 2017

b) changes in external and internal issues that are relevant to the quality management system;

c) information on the performance and effectiveness of the quality management system, including trends in:

1) customer satisfaction and feedback from relevant interested parties;

2) the extent to which quality objectives have been met;

3) process performance and conformity of products and services;

4) nonconformities and corrective actions;

5) monitoring and measurement results;

6) audit results;

7) the performance of external providers;

d) the adequacy of resources;

e) the effectiveness of actions taken to address risks and opportunities (see 6.1);

f) opportunities for improvement.

Management review inputs

Nov 2021

Audit 18 Management Review

Revision Document ID73320

****Date Revision 26 Oct 2021 Reviewed 26 Oct 2021**

Process: 7673

Check Expiry Dated Stock 09 Mar 2016

Process: 7671

Humanmed Non Conformances 09 Mar 2016

Process: 7427

Responsibility Allocation : VST Customer Complaints 09 Mar 2016

Process: 7391

Responsibility Allocation : VST Stock Meeting Customer Complaints Review ****Mandatory**** 09 Mar 2016

Process: 7389

Responsibility Allocation : VST Stock Meeting Returns Overview - From Customers 09 Mar 2016

Process: 7843

Review VST Product Feedback Negative 23 Sep 2017

Process: 7841

Review VST Feedback - Customer Complaints 23 Sep 2017

Process: 7840

Review VST Feedback - Customer Feedback Negative 23 Sep 2017

Process: 7862

Review The Audit Calender Screen 04 Oct 2017

Process: 7834

Financial Review 20 Sep 2017

Process: 5877

Review Company Data 17 Feb 2016

Process: 7070

Management Review 09 Mar 2016

Process: 7713

Review Roles And Responsibilitys 17 Aug 2016

Process: 7846

ISO System Management Review Viamed 26 Sep 2017

Process: 7848

Review ISO Scopes 27 Sep 2017

Process: 7849

Review Product Failures New Codes 28 Sep 2017

Process: 7871

Review Exclusion From Viamed 13485:2016 And VST 9001:2015 15 Oct 2017

Process: 7876

Maintain Update Of ISO Route Maps 21 Oct 2017

		Process: 7878 Review Possible Upcoming Regulation Changes 22 Oct 2017 Process: 7125 Responsibility Allocation : Intrastats Urgent Problems 09 Mar 2016 Process: 28 Supplier Review 16 Feb 2016 Process: 5887 Review ISO/EN Documents 24 Feb 2016 Process: 7199 Non Conformities Review Viamed 09 Mar 2016 Process: 7743 Customer Complaints Paper File 26 Sep 2016 Process: 7827 Review The Quality Policy VST 16 Sep 2017 Process: 7833 Importance Of Effective Quality Management 20 Sep 2017 Process: 6829 Supplier Review - Outstanding orders 09 Mar 2016 Process: 6832 Supplier Review Future orders 09 Mar 2016 Process: 7753 Management Meeting Warehouse 22 Nov 2016 Process: 5881 Training Records Review 18 Feb 2016 Process: 6851 Review Accident Book 09 Mar 2016 Process: 7847 Health And Safety Review 26 Sep 2017
9.3.3 The outputs of the management review shall include decisions and actions related to: a) opportunities for improvement; b) any need for changes to the quality management system; c) resource needs. The organization shall retain documented information as evidence of the results of	Audit 18 Management Review Revision Document ID73320 **Date Revision 26 Oct 2021 Reviewed 26 Oct 2021 Audit 20 Process verification to Managment Revision Document ID73324 Date Revision 26 Oct 2021 Reviewed 26 Oct 2021	

management reviews. Management review outputs		
1 Improvement		
10 Improvement		Process: 7433 Responsibility Allocation : VST Board Directors Meeting 09 Mar 2016
10.1 The organization shall determine and select opportunities for improvement and implement any necessary actions to meet customer requirements and enhance customer satisfaction. These shall include: a) improving products and services to meet requirements as well as to address future needs and expectations; b) correcting, preventing or reducing undesired effects; c) improving the performance and effectiveness of the quality management system. NOTE Examples of improvement can include correction, corrective action, continual improvement, breakthrough change, innovation and re-organization. General	Top Level Document: VOP 10 Non Conformance, Corrective and Preventive Actions Revision Document ID46915 Date Revision 02 Nov 2020 Reviewed 02 Nov 2020 Audit 14 Complaints and Corrective Actions Revision Document ID76091 **Date Revision 25 Nov 2021 Reviewed 25 Nov 2021 Chart 08 Correction and Prevention Revision Document ID8682 Date Revision 12 Oct 2011 Reviewed 12 Oct 2011 VM3COP27.09 Reduce goldmine Mailbox preventative maintenance Revision Document ID14907 Date Revision 02 Apr 2015 Reviewed 02 Apr 2015	Process: 7825 Responsibility Allocation : Order Picking 06 Sep 2017 Process: 7822 Review Oxylink Stock 26 Jul 2017 Process: 7387 Responsibility Allocation : VST Stock Meeting Purchase Order Requirements 09 Mar 2016
10.2 Nonconformity and corrective action	Top Level Document: VOP 10 Non Conformance, Corrective and Preventive Actions Revision Document ID46915 Date Revision 02 Nov 2020 Reviewed 02 Nov 2020	Process: 7671 Humanmed Non Conformances 09 Mar 2016
10.2.1	Top Level Document: VOP 10 Non	Process: 7830

When a nonconformity occurs, including any arising from complaints, the organization shall: a) react to the nonconformity and, as applicable: 1) take action to control and correct it; 2) deal with the consequences; b) evaluate the need for action to eliminate the cause(s) of the nonconformity, in order that it does not recur or occur elsewhere, by: 1) reviewing and analysing the nonconformity; 2) determining the causes of the nonconformity; 3) determining if similar nonconformities exist, or could potentially occur; c) implement any action needed; d) review the effectiveness of any corrective action taken; e) update risks and opportunities determined during planning, if necessary; f) make changes to the quality management system, if necessary. Corrective actions shall be appropriate to the effects of the nonconformities encountered.	Conformance, Corrective and Preventive Actions Revision Document ID46915 Date Revision 02 Nov 2020 Reviewed 02 Nov 2020 Top Level Document: VOP 19 FeedBack Customer Complaints Vigilance and Notifications VST Ltd Revision Document ID75995 **Date Revision 24 Nov 2021 Reviewed 24 Nov 2021 Audit 10 Documentation Control Revision Document ID63807 Date Revision 30 Jun 2021 Reviewed 30 Jun 2021 Audit 12 CE Files Revision Document ID63815 Date Revision 30 Jun 2021 Reviewed 30 Jun 2021 Audit 14 Complaints and Corrective Actions Revision Document ID76091 **Date Revision 25 Nov 2021 Reviewed 25 Nov 2021	Review Q.A. Failures Report 18 Sep 2017 Process: 7748 Check Repair Orders 10 Oct 2016 Process: 7427 Responsibility Allocation : VST Customer Complaints 09 Mar 2016 Process: 7391 Responsibility Allocation : VST Stock Meeting Customer Complaints Review **Mandatory** 09 Mar 2016 Process: 7841 Review VST Feedback - Customer Complaints 23 Sep 2017
10.2.2 The organization shall retain documented information as evidence of: a) the nature of the nonconformities and any subsequent actions taken; b) the results of any corrective action.	Top Level Document: VOP 19 FeedBack Customer Complaints Vigilance and Notifications Viamed Ltd Revision Document ID75475 Date Revision 18 Nov 2021 Reviewed 18 Nov 2021 Top Level Document: VOP 19 FeedBack Customer Complaints Vigilance and Notifications VST Ltd Revision Document ID75995 **Date Revision 24 Nov 2021 Reviewed 24 Nov 2021	

	Top Level Document: VOP 10 Non Conformance, Corrective and Preventive Actions Revision Document ID46915 Date Revision 02 Nov 2020 Reviewed 02 Nov 2020 Audit 10 Documentation Control Revision Document ID63807 Date Revision 30 Jun 2021 Reviewed 30 Jun 2021	
10.3 The organization shall continually improve the suitability, adequacy and effectiveness of the quality management system. The organization shall consider the results of analysis and evaluation, and the outputs from management review, to determine if there are needs or opportunities that shall be addressed as part of continual improvement. Continual improvement	Audit 10 Documentation Control Revision Document ID63807 Date Revision 30 Jun 2021 Reviewed 30 Jun 2021 Audit 18 Management Review Revision Document ID73320 **Date Revision 26 Oct 2021 Reviewed 26 Oct 2021	

Document ID	Sub Processes
ID24442	VST ISO 9001:2015 Scope Process: 7848 Review ISO Scopes 27 Sep 2017
ID73320	Audit 18 Management Review Process: 55 Business Continuity Plan 17 Feb 2016 Process: 23 Company Objectives 16 Feb 2016 Process: 6813 Management Meeting Turnover Report 09 Mar 2016 Process: 27 Management Reviews And Quality Audits 16 Feb 2016 Process: 22 Company Policys 16 Feb 2016 Process: 7750 Meeting With Management 14 Oct 2016 Process: 7793 Team Review Meeting 16 Mar 2017 Process: 7753 Management Meeting Warehouse 22 Nov 2016 Process: 6861 Management Meeting Review Weekly Meeting 09 Mar 2016

Process: 7833 Importance Of Effective Quality Management 20 Sep 2017
 Process: 7834 Financial Review 20 Sep 2017
 Process: 26 Company Resources 16 Feb 2016
 Process: 30 Responsibility Allocation : MHRA Licences And Notifications 16 Feb 2016
 Process: 31 Responsibility Allocation : Notified Body Notifications 16 Feb 2016
 Process: 32 MDALL Listings 16 Feb 2016
 Process: 7057 Responsibility Allocation : Complaints and Vigilance Notifications 09 Mar 2016
 Process: 7070 Management Review 09 Mar 2016
 Process: 29 Responsibility Allocation : CMDCAS Updates And Licences 16 Feb 2016
 Process: 5889 Responsibility Allocation : Audit And Task - Audit 24 Feb 2016
 Process: 7744 FDA Device Establishment Registration And Listing 28 Sep 2016
 Process: 7829
 Process: 6871 ISO14001 Environmental management systems 09 Mar 2016
 Process: 7874 Review For Latest Version Med Dev 2.12. 18 Oct 2017
 Process: 7877 Disaster Planning 21 Oct 2017
 Process: 7876 Maintain Update Of ISO Route Maps 21 Oct 2017
 Process: 7878 Review Possible Upcoming Regulation Changes 22 Oct 2017
 Process: 7886 Audit 18 Management Review Viamed 24 Oct 2017
 Process: 7887 Audit 18 Management Review VST 24 Oct 2017
 Process: 7890 New UPS Rates Needs Checking 24 Oct 2017
 Process: 7888 Review Processes Linked To VOPs And Audits 24 Oct 2017
 Process: 7895 FDA Device Establishment Registration 29 Oct 2017
 Process: 7912 Review The Personel Information We Collect Or Store 20 Sep 2018
 Process: 7913 Review Personnel Files 20 Sep 2018
 Process: 7918 Backup Jeans Local Folder 08 Nov 2018
 Process: 7964 Check Roles And Tasks For Incomplete Data 29 Oct 2020
 Process: 7980 Review Gov Website For Applicable Required Standards ISO9001 15 Nov 2021
 Process: 7977 Review The Agenda For The Management Review / Board Meeting Prior To The Annual Meeting 11 Nov 2021
 Process: 7978 Regulatory Requirements 11 Nov 2021
 Process: 7979 Review The Template Of The QC 21 Form To Ensure It Is Current And Valid 12 Nov 2021
 Process: 7981 ****Review Process Updates For Risk To Systems 07 Dec 2021**

ID75407

VOP 01 Documentation and Records, Control, Creation, Storage, Retrieval, Revision Control and Online Records

Process: 5940 Thumb Nail Processor 07 Mar 2016
 Process: 7827 Review The Quality Policy VST 16 Sep 2017
 Process: 7828 Review The Quality Policy Viamed 16 Sep 2017
 Process: 5934 Responsibility Allocation : Staff Training 05 Mar 2016
 Process: 7032 Responsibility Allocation : Document Requirements 09 Mar 2016
 Process: 41 Responsibility Allocation : Documentation Control 16 Feb 2016
 Process: 59 Out Of Date Documents 17 Feb 2016

Process: 5851 Duplicate Documents 17 Feb 2016
Process: 5852 Responsibility Allocation : Retention Of Records 17 Feb 2016
Process: 7130 Intrastats Information for Intrastats and L Drive 09 Mar 2016
Process: 5890 Check Website ISO Documents 24 Feb 2016
Process: 7200 Responsibility Allocation : ISO Issues 09 Mar 2016
Process: 7744 FDA Device Establishment Registration And Listing 28 Sep 2016
Process: 7941 Check Leaflets, Letterhead And Other Paperwork To See If The Correct BSI Logo Is In Use. Remove All Old If Found. 23 Sep 2019

ID63807

Audit 10 Documentation Control

Process: 10 Distribution Of Emails 16 Feb 2016
Process: 5939 Responsibility Allocation : Email ISP Routing 05 Mar 2016
Process: 5940 Thumb Nail Processor 07 Mar 2016
Process: 11 Distribution Of Mail 16 Feb 2016
Process: 6 Responsibility Allocation : Updating Contact Management System 16 Feb 2016
Process: 52 Software Verification Clear Down Backup Emails 16 Feb 2016
Process: 53 Emails 16 Feb 2016
Process: 7672 Off Site Backup 09 Mar 2016
Process: 7700 Domain Name Management 19 May 2016
Process: 9 Distribution Of Faxes 16 Feb 2016
Process: 15 Filing and Archiving 16 Feb 2016
Process: 7711 Import Bank CSV 01 Jul 2016
Process: 7722 Audit 10 Documentation Control Viamed 24 Aug 2016
Process: 7693 Collect Repair Filing From Warehouse 22 Apr 2016
Process: 12 Responsibility Allocation : Sales And Technical Information Processing 16 Feb 2016
Process: 16 Responsibility Allocation : Photocopying 16 Feb 2016
Process: 5901 Link Call Log Contacts To The CRM 02 Mar 2016
Process: 7699 Shred Sensitive Paperwork In JL Office 19 May 2016
Process: 7705 Checking For Uploaded Files 08 Jun 2016
Process: 7754
Process: 7770 Audit 10 Documentation Control VST 08 Feb 2017
Process: 6938 Responsibility Allocation : Customer Database Updates 09 Mar 2016
Process: 6940 Responsibility Allocation : Customer Ongoing task List 09 Mar 2016
Process: 7090 Responsibility Allocation : Office Procedures 09 Mar 2016
Process: 7032 Responsibility Allocation : Document Requirements 09 Mar 2016
Process: 41 Responsibility Allocation : Documentation Control 16 Feb 2016
Process: 59 Out Of Date Documents 17 Feb 2016
Process: 5851 Duplicate Documents 17 Feb 2016
Process: 5852 Responsibility Allocation : Retention Of Records 17 Feb 2016
Process: 7124 Responsibility Allocation : Intrastats 09 Mar 2016

Process: 7125 Responsibility Allocation : Intrastats Urgent Problems 09 Mar 2016
Process: 7126 Intrastats Requested Page updates 09 Mar 2016
Process: 7127 Responsibility Allocation : Intrastats Unfinished in progress Processes 09 Mar 2016
Process: 7128 Responsibility Allocation : Intrastats Future Features needed 09 Mar 2016
Process: 7129 Intrastats Cross Reference Database Tables Updates 09 Mar 2016
Process: 7130 Intrastats Information for Intrastats and L Drive 09 Mar 2016
Process: 7131 Responsibility Allocation : Intrastats Opera 09 Mar 2016
Process: 7133 Responsibility Allocation : Intrastats Contact Manager 09 Mar 2016
Process: 7739 Intrastats Amendment Log 12 Sep 2016
Process: 5877 Review Company Data 17 Feb 2016
Process: 44 Secure Socket Level Certificate 16 Feb 2016
Process: 5890 Check Website ISO Documents 24 Feb 2016
Process: 7863 Maintain Repair Codes List 05 Oct 2017
Process: 7922 Back Up Emily's Accounts Docs 04 Jan 2019

ID73324

Audit 20 Process verification to Managment

Process: 7701 AWS Amazon Web Services 23 May 2016
Process: 7723 Audit 10b Process Verification Viamed 24 Aug 2016
Process: 7730 Audit 20 Process Verification To Managment Viamed 24 Aug 2016
Process: 7827 Review The Quality Policy VST 16 Sep 2017
Process: 7828 Review The Quality Policy Viamed 16 Sep 2017
Process: 7771 Audit 10b Process Verification VST 08 Feb 2017
Process: 7778 Audit 20 Process Verification To Managment VST 08 Feb 2017
Process: 6866 Internal Process Verification Complete Systems Review 09 Mar 2016
Process: 7755 Fast Hosts Invoice 08 Dec 2016
Process: 7845 7.1.4 Environment Of Operations 25 Sep 2017
Process: 7846 ISO System Management Review Viamed 26 Sep 2017
Process: 7837 Review External Parties Influencing The QMS VST / Viamed 23 Sep 2017
Process: 7832 Cleardown Emailed Invoices 20 Sep 2017
Process: 7848 Review ISO Scopes 27 Sep 2017
Process: 7851 ****Software Validation Scan Un-QA Product To Order 23 Nov 2021**
Process: 7852 Software Validation Expired Stock 01 Oct 2017
Process: 7853 Software Validation Non Sell Able Shelf 01 Oct 2017
Process: 7854 Software Validation In Production List 01 Oct 2017
Process: 7855 Software Validation - Production Lists 01 Oct 2017
Process: 7856 Software Validation Unchecked Orders 01 Oct 2017
Process: 7857 Software Validation Stock Tracking Check 01 Oct 2017
Process: 7858 Software Validation Attempt To QA Some Stock 01 Oct 2017
Process: 7861 Software Validation Of Training Documents Forced Reading 03 Oct 2017
Process: 7850 Software Validation Scan In Correct Product 01 Oct 2017

	Process: 7871 Review Exclusion From Viamed 13485:2016 And VST 9001:2015 15 Oct 2017 Process: 7865 Software Validation Conflicting Audits 07 Oct 2017 Process: 7870 Software Validation Non Conformance Product Risk Feedback Loop 15 Oct 2017 Process: 7879 Software Validation Scheduled Tasks And Audits 22 Oct 2017 Process: 7875 Software Validation Document Control 20 Oct 2017 Process: 7880 Software Validation Out Of Date Documents 22 Oct 2017 Process: 7881 Software Validation - Live Orders 22 Oct 2017
ID8700	Chart 27 Customer Complaints Chart 27 Process: 7743 Customer Complaints Paper File 26 Sep 2016
ID70147	Audit 08 Training, Competence and Human Resources Process: 7720 Audit 08 Training Viamed 24 Aug 2016 Process: 6839 Responsibility Allocation : Personnel Holidays and Time Adjustments 09 Mar 2016 Process: 5881 Training Records Review 18 Feb 2016 Process: 5904 Responsibility Allocation : Taking On New Staff 02 Mar 2016 Process: 5936 Wages Calculations 05 Mar 2016 Process: 6837 Personnel Requirements and Training 09 Mar 2016 Process: 6851 Review Accident Book 09 Mar 2016 Process: 6877 Responsibility Allocation : Alarm Key Holders 09 Mar 2016 Process: 6906 Responsibility Allocation : Time Working Away 09 Mar 2016 Process: 6928 Responsibility Allocation : Staff 09 Mar 2016 Process: 7074 Process: 7759 Health Declaration Sheet 23 Jan 2017 Process: 7768 Audit 08 Training VST 08 Feb 2017 Process: 5934 Responsibility Allocation : Staff Training 05 Mar 2016 Process: 6841 Responsibility Allocation : Grants 09 Mar 2016 Process: 7070 Management Review 09 Mar 2016 Process: 7713 Review Roles And Responsibilitys 17 Aug 2016 Process: 7883 Appraisal 23 Oct 2017 Process: 7884 Pay Review 23 Oct 2017 Process: 7908 Private Information Data 27 Jul 2018 Process: 7907 Annual Review Doc Management 27 Jul 2018 Process: 7937 Diversity Impact Assessment 27 Jun 2019 Process: 7951 Server Review 05 Mar 2020 Process: 7982 Check There Are No Changes To Employment Law 21 Nov 2021 Process: 7983 To Check On Line And See If There Have Been Any Changes To Gdpr We Need To Be Aware Of. 21 Nov 2021
ID22684	VM3COP00.00 Viamed Quality Statement policy and objectives Process: 23 Company Objectives 16 Feb 2016 Process: 22 Company Policys 16 Feb 2016

	Process: 7828 Review The Quality Policy Viamed 16 Sep 2017 Process: 7833 Importance Of Effective Quality Management 20 Sep 2017
ID22062	VM3COP00.00 VST Quality Statement policy and objectives Process: 23 Company Objectives 16 Feb 2016 Process: 7827 Review The Quality Policy VST 16 Sep 2017 Process: 7833 Importance Of Effective Quality Management 20 Sep 2017
ID27474	VM3COP02.02 Viamed Company Responsibility's organisation chart structure Process: 5877 Review Company Data 17 Feb 2016
ID73529	VOP 02 Personnel and Responsibility , Staff and Staffing Issues, Training, Roles and Tasks Process: 39 Environmental Policy Document Review 16 Feb 2016 Process: 7741 Review Ethical Policy 14 Sep 2016 Process: 6839 Responsibility Allocation : Personnel Holidays and Time Adjustments 09 Mar 2016 Process: 5881 Training Records Review 18 Feb 2016 Process: 5904 Responsibility Allocation : Taking On New Staff 02 Mar 2016 Process: 6837 Personnel Requirements and Training 09 Mar 2016 Process: 6877 Responsibility Allocation : Alarm Key Holders 09 Mar 2016 Process: 6906 Responsibility Allocation : Time Working Away 09 Mar 2016 Process: 6928 Responsibility Allocation : Staff 09 Mar 2016 Process: 7074 Process: 7042 Responsibility Allocation : Work Environment 09 Mar 2016 Process: 5934 Responsibility Allocation : Staff Training 05 Mar 2016 Process: 5874 Childcare Vouchers Edenred 17 Feb 2016 Process: 7753 Management Meeting Warehouse 22 Nov 2016 Process: 34 Responsibility Allocation : Insurance Is Upto Date 16 Feb 2016 Process: 5869 Responsibility Allocation : Legal Company Car Registration 17 Feb 2016 Process: 6841 Responsibility Allocation : Grants 09 Mar 2016 Process: 6843 Process: 6861 Management Meeting Review Weekly Meeting 09 Mar 2016 Process: 30 Responsibility Allocation : MHRA Licences And Notifications 16 Feb 2016 Process: 31 Responsibility Allocation : Notified Body Notifications 16 Feb 2016 Process: 32 MDALL Listings 16 Feb 2016 Process: 7033 Responsibility Allocation : Management commitment to ISO 09 Mar 2016 Process: 7037 Responsibility Allocation : Responsibility, authority and communication 09 Mar 2016 Process: 7057 Responsibility Allocation : Complaints and Vigilance Notifications 09 Mar 2016 Process: 7713 Review Roles And Responsibility's 17 Aug 2016 Process: 7837 Review External Parties Influencing The QMS VST / Viamed 23 Sep 2017 Process: 29 Responsibility Allocation : CMDCAS Updates And Licences 16 Feb 2016 Process: 7848 Review ISO Scopes 27 Sep 2017

	Process: 7891 Fire Alarm Evacuation Drill 25 Oct 2017 Process: 7908 Private Information Data 27 Jul 2018 Process: 7907 Annual Review Doc Management 27 Jul 2018 Process: 7937 Diversity Impact Assessment 27 Jun 2019 Process: 7961 R D Room - Tidy, Empty Bins, Remove Cups. Caution Around Oxygen Supply 05 Oct 2020 Process: 7982 Check There Are No Changes To Employment Law 21 Nov 2021 Process: 7983 To Check On Line And See If There Have Been Any Changes To Gdpr We Need To Be Aware Of. 21 Nov 2021
ID17423	VM3COP02 Organisation Responsibilities Viamed Process: 6967 Responsibility Allocation : VIAMED Stock Meeting Repairs Review - Pulse Oximetry Sensors 09 Mar 2016 Process: 7900 Royal Mail - Mail Retention Form 29 Mar 2018
ID21800	VM3COP19 Health and Safety Process: 6855 Risk Assessment HSE 09 Mar 2016
ID22429	Viamed Top Level Quality Objectives Process: 23 Company Objectives 16 Feb 2016
ID31036	VOP 18 Maintenance Building, Fabric and Infrastructure Process: 5856 Cleaning The Kitchen 17 Feb 2016 Process: 5853 Vacuuming Of The Office, Hall And Meeting Room 17 Feb 2016 Process: 5900 Cleaning Of Office Windows 25 Feb 2016 Process: 5878 Empty Office Bins 18 Feb 2016 Process: 5912 Responsibility Allocation : Main Recycle Bins 03 Mar 2016 Process: 5906 Empty Paper Bins 03 Mar 2016 Process: 7805 Empty Kitchen Bins 22 May 2017 Process: 5909 Empty Warehouse Bins 03 Mar 2016 Process: 7706 Update Virus Software And Scan For Viruses 10 Jun 2016 Process: 7802 Clean Kitchen Sides 22 May 2017 Process: 7803 Dishwashing 22 May 2017 Process: 7804 Sweep Kitchen Floor 22 May 2017 Process: 7806 Watering Plants 22 May 2017 Process: 7807 Process: 54 Responsibility Allocation : Gents Toilets 17 Feb 2016 Process: 5907 Hoover Warehouse 03 Mar 2016 Process: 5908 Sweep Warehouse 03 Mar 2016 Process: 5910 Clean Duckets 03 Mar 2016 Process: 5911 Clear Cardboard 03 Mar 2016 Process: 7698 Clean Toilets 17 May 2016 Process: 7131 Responsibility Allocation : Intrastats Opera 09 Mar 2016 Process: 7133 Responsibility Allocation : Intrastats Contact Manager 09 Mar 2016

	Process: 7132 Responsibility Allocation : Intrastats Goldmine 09 Mar 2016 Process: 7896 Tree In Car Park 22 Dec 2017
ID69457	Audit 16 Sales and Marketing Process: 21 Office Sales Projects 16 Feb 2016 Process: 17 Process: 40 Responsibility Allocation : Calender 16 Feb 2016 Process: 5870 Book Arab Health 17 Feb 2016 Process: 19 Maintaining Leaflet Stocks 16 Feb 2016 Process: 20 Processing Of Mail Shots 16 Feb 2016 Process: 5873 Distributor Contract Reviews 17 Feb 2016 Process: 5885 Responsibility Allocation : Monthly Reports 24 Feb 2016 Process: 5883 Responsibility Allocation : Monthly Sales Report 24 Feb 2016 Process: 6888 Viamed Automotive UK 09 Mar 2016 Process: 6898 GHX Web Pricing 09 Mar 2016 Process: 5884 Responsibility Allocation : Monthly Report 24 Feb 2016 Process: 5886 Responsibility Allocation : Monthly Report 24 Feb 2016 Process: 6891 Responsibility Allocation : Exhibitions Co-ordinator 09 Mar 2016 Process: 7909 EAN GTIN Online Database 06 Aug 2018 Process: 7920 Sales Warnings 20 Dec 2018 Process: 7927 Contract Pricing Review 14 Feb 2019 Process: 7926 Sales Forecasts Export 22 Jan 2019 Process: 7921 VST Bags And Grey Sensor 03 Jan 2019 Process: 7925 Providing Ebay Feedback 16 Jan 2019 Process: 7916 Google Webmaster Tools 16 Oct 2018 Process: 7931 Competitor Pricing 14 Mar 2019 Process: 7949 Sales Projects Send To Sales Team 04 Mar 2020 Process: 7947 8010004 - JJ-CCR Oxygen Sensor Orders 04 Mar 2020 Process: 7948 8010006 - REVo Oxygen Sensor Orders 04 Mar 2020 Process: 7950 Envitec Oxygen Sensor Parts Stock Check 05 Mar 2020 Process: 7959 Audit 16 Sales And Marketing Viamed 28 Sep 2020 Process: 7960 Audit 16 Sales And Marketing VST 28 Sep 2020
ID69328	Audit 02 Contract Review and Sales Order Processing Process: 5 Responsibility Allocation : Processing Of Sales Orders 16 Feb 2016 Process: 36 Emailing Of Invoices 16 Feb 2016 Process: 5892 Checking EBay And Amazon For Orders And Messages 25 Feb 2016 Process: 5894 Checking Of Active List 25 Feb 2016 Process: 7 Responsibility Allocation : Checking Of Sales Orders 16 Feb 2016 Process: 5943 Check Cardea And Multiquote 08 Mar 2016 Process: 5891 Processing Of Repair Quotes And Orders 25 Feb 2016

Process: 2 Answering Telephones 16 Feb 2016
Process: 37 West Yorkshire Ambulance Stock 16 Feb 2016
Process: 5945 Responsibility Allocation : Sending Samples 08 Mar 2016
Process: 5946 Responsibility Allocation : Sending Sale Or Returns 08 Mar 2016
Process: 5948 Adding New Accounts To Opera 08 Mar 2016
Process: 5949 Filling Credit Card Slips 08 Mar 2016
Process: 5895 Responsibility Allocation : Completing Office Job List 25 Feb 2016
Process: 5875 Check Paypal For Orders 17 Feb 2016
Process: 7675 Responsibility Allocation : Ordering Demo Stock For Humanmed Reps 11 Mar 2016
Process: 5944 Responsibility Allocation : Chasing Lost Customers 08 Mar 2016
Process: 3 Responsibility Allocation : Meeting And Greeting Visitors To The Company 16 Feb 2016
Process: 4 Responsibility Allocation : Assisting With Refreshments For Visitors 16 Feb 2016
Process: 7676 PDFing Of Invoices Viamed 17 Mar 2016
Process: 7696 Send VIAMED Delivery Notifications 28 Apr 2016
Process: 5893 Answering Website Questions 25 Feb 2016
Process: 7678 Check Catalog 360 Circle For Quotes And Orders 08 Apr 2016
Process: 5899 Proforma And Quote Chasing 25 Feb 2016
Process: 7710 Responsibility Allocation : Proforma And Quote Processing 29 Jun 2016
Process: 14 Fax Paper 16 Feb 2016
Process: 5882 Responsibility Allocation : Send Post To Humanmed 24 Feb 2016
Process: 7715 Audit 02 Contract Review Viamed 24 Aug 2016
Process: 7734 Responsibility Allocation : Humanmed Order Processing 25 Aug 2016
Process: 7677
Process: 5897 Responsibility Allocation : Franking Mail 25 Feb 2016
Process: 7709 Delivered not Invoiced 28 Jun 2016
Process: 6954 Back Orders Review - By Customer 09 Mar 2016
Process: 8 Responsibility Allocation : Order And Status Liaison With Customers 16 Feb 2016
Process: 5896 Responsibility Allocation : Ensuring ORD's Are Taken To Goods Out And Invoices Are Retrieved 25 Feb 2016
Process: 5913 Check For Humanmed Orders In Logistics Mailbox 03 Mar 2016
Process: 5947 Responsibility Allocation : Search For Distributors 08 Mar 2016
Process: 6958 Responsibility Allocation : Shipped Order Queries 09 Mar 2016
Process: 7686 Thorough Checking Of Awaiting Action Tray 21 Apr 2016
Process: 7712 Review Inward Payments 01 Jul 2016
Process: 7735 Ensure SOR's Are Followed Up 01 Sep 2016
Process: 7758 Check For GHX Orders 17 Jan 2017
Process: 7761 Send VST Delivery Notifications 01 Feb 2017
Process: 7783 PDF VST Invoices And Purchase Orders 10 Feb 2017
Process: 7795 Answering UK Web Questions 27 Apr 2017
Process: 7822 Review Oxylink Stock 26 Jul 2017

	Process: 7791 Price List Check 10 Mar 2017 Process: 7763 Audit 02 Contract Review VST 08 Feb 2017 Process: 7808 Ensure All Invoice Correctly Tagged 02 Jun 2017 Process: 5872 Check Sale Or Returns Export 17 Feb 2016 Process: 5871 Check Sale Or Returns 17 Feb 2016 Process: 5876 E.Commerce Cardea And Multiquote 17 Feb 2016 Process: 7782 Remove Started But Not Used Order Numbers 08 Feb 2017 Process: 6956 Responsibility Allocation : Sales Order Issues 09 Mar 2016 Process: 6921 Responsibility Allocation : Customer pricing agreements 09 Mar 2016 Process: 6922 Process: 6959 Responsibility Allocation : Sales Forward Orders Review 09 Mar 2016 Process: 7801 VST Price Review 17 May 2017 Process: 5905 Responsibility Allocation : Price Checking 02 Mar 2016 Process: 6950 Process: 7697 Yearly Pricing Review 09 May 2016 Process: 7670 Humanmed general Issues 09 Mar 2016 Process: 7872 Embargo Countries NOT Allowed To Sell To 16 Oct 2017 Process: 7893 VST Price Lists 28 Oct 2017 Process: 7894 VST Customer Agreements 28 Oct 2017 Process: 7936 B2B Router / Peppol Responsibilitys 19 Jun 2019 Process: 7941 Check Leaflets, Letterhead And Other Paperwork To See If The Correct BSI Logo Is In Use. Remove All Old If Found. 23 Sep 2019 Process: 7953 Vandagraph Delivery Notifications 26 May 2020 Process: 7954 Vandagraph Email Of Invoices 26 May 2020 Process: 7955 Vandagraph Shipper SignOff Collection 26 May 2020 Process: 7970 Proforma And Quote Chasing Ryan 31 Aug 2021 Process: 7971 Proforma And Quote Chasing Steve Hardaker 31 Aug 2021
ID51629	Audit 01 Picking packing Process: 7714 Audit 01 Picking Packing Viamed 24 Aug 2016 Process: 7825 Responsibility Allocation : Order Picking 06 Sep 2017 Process: 5859 Review Un-shipped Parcels 17 Feb 2016 Process: 6970 Process: 7691 Ship Sale Or Returns 21 Apr 2016 Process: 7762 Audit 01 Picking Packing VST 08 Feb 2017 Process: 7796 Review Franking Label Errors 08 May 2017 Process: 7797 Check Order Are Being Picked In Priority Order 10 May 2017 Process: 7798 Orders And Items Shipped Per Month 10 May 2017 Process: 7860 Goods Out Picking 03 Oct 2017
ID75973	VOP 07 Stock Control, Handling, Control of Labelling, Storage, Movement

Process: 6973 Responsibility Allocation : Stock Transfers. (QC19) 09 Mar 2016
Process: 7675 Responsibility Allocation : Ordering Demo Stock For Humanmed Reps 11 Mar 2016
Process: 5872 Check Sale Or Returns Export 17 Feb 2016
Process: 5871 Check Sale Or Returns 17 Feb 2016
Process: 5855 Purchase Order Requirements Teledyne 17 Feb 2016
Process: 5858 Opera Stock Adjustments 17 Feb 2016
Process: 5868 Return Goods To Suppliers 17 Feb 2016
Process: 5935 Stock Allocations 05 Mar 2016
Process: 6829 Supplier Review - Outstanding orders 09 Mar 2016
Process: 6832 Supplier Review Future orders 09 Mar 2016
Process: 6840
Process: 6848
Process: 6850 Current Stock Levels 09 Mar 2016
Process: 6945 Missing Stock or Adjustments 09 Mar 2016
Process: 6955 Production Requirements 09 Mar 2016
Process: 7046 Responsibility Allocation : Stock Purchasing 09 Mar 2016
Process: 7051 Responsibility Allocation : Control of nonconforming product 09 Mar 2016
Process: 7673 Check Expiry Dated Stock 09 Mar 2016
Process: 7679 Check Stock Requirements Supplier Teledyne 18 Apr 2016
Process: 7680 Check Stock Requirements Supplier Envitec 18 Apr 2016
Process: 7681 Check Stock Requirements Supplier Posey 18 Apr 2016
Process: 7682 Check Stock Requirements Supplier Bluepoint 18 Apr 2016
Process: 7687 Vandagraph Duckets 21 Apr 2016
Process: 7688
Process: 7689 Move Stock From QA Shelf To Stock Shelf Monday 21 Apr 2016
Process: 7694 Move Stock From QA Shelf To Stock Shelf Tuesday 28 Apr 2016
Process: 7695 Top Up Quick Shipping Shelves 28 Apr 2016
Process: 7708 Acorn 0014904 17 Jun 2016
Process: 7798 Orders And Items Shipped Per Month 10 May 2017
Process: 6961 Responsibility Allocation : VIAMED Stock Meeting Purchase Order Requirements 09 Mar 2016
Process: 7683 Check Stock For Proforma 18 Apr 2016
Process: 6968 Responsibility Allocation : VIAMED Stock Meeting Repairs Review - General 09 Mar 2016
Process: 6949 Responsibility Allocation : VIAMED Stock Meeting QA Processing 09 Mar 2016
Process: 6948 Responsibility Allocation : VIAMED Stock Meeting Stock Processing 09 Mar 2016
Process: 6947 Responsibility Allocation : VIAMED Stock Meeting Stock Queries 09 Mar 2016
Process: 7830 Review Q.A. Failures Report 18 Sep 2017
Process: 7864 ESD Work Stations 07 Oct 2017
Process: 7873 On Site Environment Review 18 Oct 2017
Process: 7866 Oxygen Cylinder Check 13 Oct 2017

	Process: 7897 Daily O2 Sensors Returns 04 Jan 2018 Process: 7909 EAN GTIN Online Database 06 Aug 2018 Process: 7943 Review Stocks Of 8000004 01 Oct 2019 Process: 7944 Sealant, Glues, Greases, Sprays, Gases And Tapes You Use In Production, Service And Repairs For Viamed And VST 09 Oct 2019 Process: 7962 VST Supplier QA Results 28 Oct 2020 Process: 7967 VST Stock Count For End April 01 Jul 2021 Process: 7969 Weee Waste Reporting 23 Aug 2021
ID75475	VOP 19 Feedback Customer Complaints Vigilance and Notifications Viamed Ltd Process: 7743 Customer Complaints Paper File 26 Sep 2016 Process: 7671 Humanmed Non Conformances 09 Mar 2016 Process: 6931 Customer Complaints 09 Mar 2016 Process: 7839 Review VIAMED Feedback - Customer Complaints 23 Sep 2017 Process: 7838 Review VIAMED Feedback - Customer Feedback Negative 23 Sep 2017 Process: 7070 Management Review 09 Mar 2016 Process: 7840 Review VST Feedback - Customer Feedback Negative 23 Sep 2017 Process: 7841 Review VST Feedback - Customer Complaints 23 Sep 2017 Process: 7842 Review VIAMED Product Feedback Negative 23 Sep 2017 Process: 7843 Review VST Product Feedback Negative 23 Sep 2017 Process: 7174 Process: 7175 Process: 7179 Process: 7874 Review For Latest Version Med Dev 2.12. 18 Oct 2017 Process: 7979 Review The Template Of The QC 21 Form To Ensure It Is Current And Valid 12 Nov 2021
ID18641	VM3COP20.01 Post In Distributing the Post Process: 11 Distribution Of Mail 16 Feb 2016 Process: 5882 Responsibility Allocation : Send Post To Humanmed 24 Feb 2016
ID75949	VOP 03 Contract Review, Enquires, Office Processes Process: 5 Responsibility Allocation : Processing Of Sales Orders 16 Feb 2016 Process: 10 Distribution Of Emails 16 Feb 2016 Process: 36 Emailing Of Invoices 16 Feb 2016 Process: 5892 Checking EBay And Amazon For Orders And Messages 25 Feb 2016 Process: 5894 Checking Of Active List 25 Feb 2016 Process: 7 Responsibility Allocation : Checking Of Sales Orders 16 Feb 2016 Process: 5943 Check Cardea And Multiquote 08 Mar 2016 Process: 5891 Processing Of Repair Quotes And Orders 25 Feb 2016 Process: 11 Distribution Of Mail 16 Feb 2016 Process: 2 Answering Telephones 16 Feb 2016

Process: 37 West Yorkshire Ambulance Stock 16 Feb 2016
Process: 5948 Adding New Accounts To Opera 08 Mar 2016
Process: 5949 Filling Credit Card Slips 08 Mar 2016
Process: 6 Responsibility Allocation : Updating Contact Management System 16 Feb 2016
Process: 5895 Responsibility Allocation : Completing Office Job List 25 Feb 2016
Process: 5875 Check Paypal For Orders 17 Feb 2016
Process: 5944 Responsibility Allocation : Chasing Lost Customers 08 Mar 2016
Process: 3 Responsibility Allocation : Meeting And Greeting Visitors To The Company 16 Feb 2016
Process: 4 Responsibility Allocation : Assisting With Refreshments For Visitors 16 Feb 2016
Process: 7676 PDFing Of Invoices Viamed 17 Mar 2016
Process: 9 Distribution Of Faxes 16 Feb 2016
Process: 7696 Send VIAMED Delivery Notifications 28 Apr 2016
Process: 5857 Customer Service Logs 17 Feb 2016
Process: 5893 Answering Website Questions 25 Feb 2016
Process: 7678 Check Catalog 360 Circle For Quotes And Orders 08 Apr 2016
Process: 15 Filing and Archiving 16 Feb 2016
Process: 5899 Proforma And Quote Chasing 25 Feb 2016
Process: 7710 Responsibility Allocation : Proforma And Quote Processing 29 Jun 2016
Process: 7707 Send Purchase Orders To Suppliers 13 Jun 2016
Process: 14 Fax Paper 16 Feb 2016
Process: 5882 Responsibility Allocation : Send Post To Humanmed 24 Feb 2016
Process: 7734 Responsibility Allocation : Humanmed Order Processing 25 Aug 2016
Process: 5850 Purchase Order Log 17 Feb 2016
Process: 7693 Collect Repair Filing From Warehouse 22 Apr 2016
Process: 7677
Process: 5897 Responsibility Allocation : Franking Mail 25 Feb 2016
Process: 21 Office Sales Projects 16 Feb 2016
Process: 7709 Delivered not Invoiced 28 Jun 2016
Process: 8 Responsibility Allocation : Order And Status Liaison With Customers 16 Feb 2016
Process: 12 Responsibility Allocation : Sales And Technical Information Processing 16 Feb 2016
Process: 16 Responsibility Allocation : Photocopying 16 Feb 2016
Process: 17
Process: 20 Processing Of Mail Shots 16 Feb 2016
Process: 5896 Responsibility Allocation : Ensuring ORD's Are Taken To Goods Out And Invoices Are Retrieved 25 Feb 2016
Process: 5901 Link Call Log Contacts To The CRM 02 Mar 2016
Process: 5913 Check For Humanmed Orders In Logistics Mailbox 03 Mar 2016
Process: 5947 Responsibility Allocation : Search For Distributors 08 Mar 2016
Process: 6958 Responsibility Allocation : Shipped Order Queries 09 Mar 2016
Process: 7686 Thorough Checking Of Awaiting Action Tray 21 Apr 2016

Process: 7699 Shred Sensitive Paperwork In JL Office 19 May 2016
Process: 7705 Checking For Uploaded Files 08 Jun 2016
Process: 7712 Review Inward Payments 01 Jul 2016
Process: 7735 Ensure SOR's Are Followed Up 01 Sep 2016
Process: 7751 VST Purchase Order Log 02 Nov 2016
Process: 7758 Check For GHX Orders 17 Jan 2017
Process: 7760 Send Service Offers 31 Jan 2017
Process: 7761 Send VST Delivery Notifications 01 Feb 2017
Process: 7783 PDF VST Invoices And Purchase Orders 10 Feb 2017
Process: 7792 Shipped Order Success Report 13 Mar 2017
Process: 7795 Answering UK Web Questions 27 Apr 2017
Process: 7822 Review Oxylink Stock 26 Jul 2017
Process: 5876 E.Commerce Cardea And Multiquote 17 Feb 2016
Process: 5873 Distributor Contract Reviews 17 Feb 2016
Process: 5885 Responsibility Allocation : Monthly Reports 24 Feb 2016
Process: 6938 Responsibility Allocation : Customer Database Updates 09 Mar 2016
Process: 6940 Responsibility Allocation : Customer Ongoing task List 09 Mar 2016
Process: 6956 Responsibility Allocation : Sales Order Issues 09 Mar 2016
Process: 5866 UPS Shipping Fuel Surcharge 17 Feb 2016
Process: 6952 Responsibility Allocation : Lost in Shipping Claims 09 Mar 2016
Process: 6971 Responsibility Allocation : Freight Courier Cost Request 09 Mar 2016
Process: 7692 Responsibility Allocation : Take Complete Repair Paperwork To Office 22 Apr 2016
Process: 7796 Review Franking Label Errors 08 May 2017
Process: 6916 Responsibility Allocation : Service exisiting 09 Mar 2016
Process: 6917 Responsibility Allocation : Service extension 09 Mar 2016
Process: 7863 Maintain Repair Codes List 05 Oct 2017
Process: 7872 Embargo Countries NOT Allowed To Sell To 16 Oct 2017
Process: 7890 New UPS Rates Needs Checking 24 Oct 2017
Process: 7893 VST Price Lists 28 Oct 2017
Process: 7894 VST Customer Agreements 28 Oct 2017
Process: 7901 UPS Exceptions Checkup 20 Apr 2018
Process: 7957 Warehouse Requests 29 May 2020
Process: 7959 Audit 16 Sales And Marketing Viamed 28 Sep 2020
Process: 7970 Proforma And Quote Chasing Ryan 31 Aug 2021
Process: 7971 Proforma And Quote Chasing Steve Hardaker 31 Aug 2021

ID77289

Process: 7731 Audit 21 Audit Of Audit Viamed 24 Aug 2016
Process: 7779 Audit 21 Audit Of Audit VST 08 Feb 2017
Process: 38 Audits Up to Date and Confirm next years Audit schedule 16 Feb 2016

	Process: 7093 BSI Audits Calander 09 Mar 2016 Process: 7670 Humanmed general Issues 09 Mar 2016 Process: 7862 Review The Audit Calender Screen 04 Oct 2017
ID21314	Process: 6828
ID63815	Audit 12 CE Files Process: 7725 Audit 12 CE Files Viamed 24 Aug 2016 Process: 7773 Audit 12 CE Files VST 08 Feb 2017 Process: 24 Responsibility Allocation : Compliance ISO Standards 16 Feb 2016 Process: 7172 Responsibility Allocation : CE Technical Files 09 Mar 2016 Process: 7071 Post Market Surveillance 09 Mar 2016
ID75461	VOP 13 Process Monitoring, System Reviews, Audits, Management Reviews and Analysis Data Process: 55 Business Continuity Plan 17 Feb 2016 Process: 23 Company Objectives 16 Feb 2016 Process: 27 Management Reviews And Quality Audits 16 Feb 2016 Process: 7714 Audit 01 Picking Packing Viamed 24 Aug 2016 Process: 7715 Audit 02 Contract Review Viamed 24 Aug 2016 Process: 7716 Audit 03 Design Control Viamed 24 Aug 2016 Process: 7717 Audit 05 Purchasing Suppliers Viamed 24 Aug 2016 Process: 7718 Audit 06 Calibration Viamed 24 Aug 2016 Process: 7719 Audit 07 Handling And Storage Viamed 24 Aug 2016 Process: 7720 Audit 08 Training Viamed 24 Aug 2016 Process: 7721 Audit 09 Goods Inward And Product Identity Viamed 24 Aug 2016 Process: 7722 Audit 10 Documentation Control Viamed 24 Aug 2016 Process: 7723 Audit 10b Process Verification Viamed 24 Aug 2016 Process: 7724 Audit 11 Repairs And Service Viamed 24 Aug 2016 Process: 7725 Audit 12 CE Files Viamed 24 Aug 2016 Process: 7726 Audit 14 Complaints And Corrective Actions Viamed 24 Aug 2016 Process: 7727 Audit 15 Production Viamed 24 Aug 2016 Process: 7728 Audit 17 Internal Audits Viamed 24 Aug 2016 Process: 7729 Audit 19 Health And Saftey Viamed 24 Aug 2016 Process: 7730 Audit 20 Process Verification To Managment Viamed 24 Aug 2016 Process: 7731 Audit 21 Audit Of Audit Viamed 24 Aug 2016 Process: 7732 Audit 22 Post Market Survellance Viamed 24 Aug 2016 Process: 7733 Audit 23 Analysis Of Data Viamed 24 Aug 2016 Process: 6828 Process: 22 Company Policys 16 Feb 2016 Process: 7754

Process: 7762 Audit 01 Picking Packing VST 08 Feb 2017
Process: 7763 Audit 02 Contract Review VST 08 Feb 2017
Process: 7764 Audit 03 Design Control VST 08 Feb 2017
Process: 7765 Audit 05 Purchasing Suppliers VST 08 Feb 2017
Process: 7766 Audit 06 Calibration VST 08 Feb 2017
Process: 7767 Audit 07 Handling And Storage VST 08 Feb 2017
Process: 7768 Audit 08 Training VST 08 Feb 2017
Process: 7769 Audit 09 Goods Inward And Product Identity VST 08 Feb 2017
Process: 7770 Audit 10 Documentation Control VST 08 Feb 2017
Process: 7771 Audit 10b Process Verification VST 08 Feb 2017
Process: 7772 Audit 11 Repairs And Service VST 08 Feb 2017
Process: 7773 Audit 12 CE Files VST 08 Feb 2017
Process: 7774 Audit 14 Complaints And Corrective Actions VST 08 Feb 2017
Process: 7775 Audit 15 Production VST 08 Feb 2017
Process: 7776 Audit 17 Internal Audits VST 08 Feb 2017
Process: 7777 Audit 19 Health And Safety VST 08 Feb 2017
Process: 7778 Audit 20 Process Verification To Management VST 08 Feb 2017
Process: 7779 Audit 21 Audit Of Audit VST 08 Feb 2017
Process: 7780 Audit 22 Post Market Surveillance VST 08 Feb 2017
Process: 7781 Audit 23 Analysis Of Data VST 08 Feb 2017
Process: 7808 Ensure All Invoice Correctly Tagged 02 Jun 2017
Process: 6886 Responsibility Allocation : VIAMED Sales And Marketing Sales Viamed Medical Export 09 Mar 2016
Process: 6887 Responsibility Allocation : VIAMED Sales And Marketing Sales Viamed Automotive Export 09 Mar 2016
Process: 7204 Responsibility Allocation : VIAMED Board Directors Meeting Distributor Issues 09 Mar 2016
Process: 24 Responsibility Allocation : Compliance ISO Standards 16 Feb 2016
Process: 28 Supplier Review 16 Feb 2016
Process: 6865 Responsibility Allocation : Non Conformance Effectiveness 09 Mar 2016
Process: 6866 Internal Process Verification Complete Systems Review 09 Mar 2016
Process: 7172 Responsibility Allocation : CE Technical Files 09 Mar 2016
Process: 7782 Remove Started But Not Used Order Numbers 08 Feb 2017
Process: 7090 Responsibility Allocation : Office Procedures 09 Mar 2016
Process: 7138 Non Conformance Issues Any New QC21 Forms 09 Mar 2016
Process: 57 Temporary Stock Notices 17 Feb 2016
Process: 5854 Stock FAQ Admin List 17 Feb 2016
Process: 7043 Responsibility Allocation : Planning of product realization 09 Mar 2016
Process: 38 Audits Up to Date and Confirm next years Audit schedule 16 Feb 2016
Process: 5877 Review Company Data 17 Feb 2016
Process: 6904 Responsibility Allocation : Sales And Marketing Internal sales 09 Mar 2016
Process: 6944 Responsibility Allocation : Stock Meeting 09 Mar 2016

Process: 7846 ISO System Management Review Viamed 26 Sep 2017
Process: 7834 Financial Review 20 Sep 2017
Process: 26 Company Resources 16 Feb 2016
Process: 7070 Management Review 09 Mar 2016
Process: 7837 Review External Parties Influencing The QMS VST / Viamed 23 Sep 2017
Process: 5887 Review ISO/EN Documents 24 Feb 2016
Process: 5889 Responsibility Allocation : Audit And Task - Audit 24 Feb 2016
Process: 7071 Post Market Surveillance 09 Mar 2016
Process: 7093 BSI Audits Calander 09 Mar 2016
Process: 7829
Process: 7670 Humanmed general Issues 09 Mar 2016
Process: 6821 Responsibility Allocation : VIAMED Management Meeting Supplier Review 09 Mar 2016
Process: 6831 Responsibility Allocation : VIAMED Management Meeting Supplier Review - Min / Max - Re-Orders 09 Mar 2016
Process: 6833 Responsibility Allocation : VIAMED Management Meeting MDA Recalls 09 Mar 2016
Process: 6834 Responsibility Allocation : VIAMED Management Meeting Additional Purchase Orders 09 Mar 2016
Process: 6836 Responsibility Allocation : VIAMED Management Meeting Research and Development rnd 09 Mar 2016
Process: 6920 Responsibility Allocation : VIAMED Sales And Marketing Price Lists UK 09 Mar 2016
Process: 6924 Responsibility Allocation : VIAMED Sales And Marketing Price Lists Export 09 Mar 2016
Process: 6935 Responsibility Allocation : VIAMED Sales And Marketing Products to be Marketed 09 Mar 2016
Process: 6936 Responsibility Allocation : VIAMED Sales And Marketing NHS Supplies Future Technology 09 Mar 2016
Process: 6941 Responsibility Allocation : VIAMED Sales And Marketing New Potential Products 09 Mar 2016
Process: 7039 Responsibility Allocation : Provision of Resources 09 Mar 2016
Process: 7187 Responsibility Allocation : VIAMED Board Directors Meeting Profitability 09 Mar 2016
Process: 7196 Responsibility Allocation : VIAMED Board Directors Meeting Stock Levels 09 Mar 2016
Process: 6871 ISO14001 Environmental management systems 09 Mar 2016
Process: 7848 Review ISO Scopes 27 Sep 2017
Process: 7862 Review The Audit Calender Screen 04 Oct 2017
Process: 7877 Disaster Planning 21 Oct 2017
Process: 7879 Software Validation Scheduled Tasks And Audits 22 Oct 2017
Process: 7876 Maintain Update Of ISO Route Maps 21 Oct 2017
Process: 7878 Review Possible Upcoming Regulation Changes 22 Oct 2017
Process: 7885 Audit 04 Accounts and Finance 23 Oct 2017
Process: 7886 Audit 18 Management Review Viamed 24 Oct 2017
Process: 7887 Audit 18 Management Review VST 24 Oct 2017
Process: 7889 Audit 24 Servicing Viamed 24 Oct 2017
Process: 7888 Review Processes Linked To VOPs And Audits 24 Oct 2017
Process: 7965 VST Feedback 29 Oct 2020
Process: 7964 Check Roles And Tasks For Incomplete Data 29 Oct 2020
Process: 7980 Review Gov Website For Applicable Required Standards ISO9001 15 Nov 2021

	Process: 7972 ISO System Management Review Vst 26 Oct 2021 Process: 7973 VST Product Performance - Customers 27 Oct 2021 Process: 7974 VST Product Performance - Suppliers 27 Oct 2021 Process: 7977 Review The Agenda For The Management Review / Board Meeting Prior To The Annual Meeting 11 Nov 2021 Process: 7978 Regulatory Requirements 11 Nov 2021 Process: 7981 **Review Process Updates For Risk To Systems 07 Dec 2021
ID31024	VOP 12 Training Process: 7750 Meeting With Management 14 Oct 2016 Process: 7793 Team Review Meeting 16 Mar 2017 Process: 5934 Responsibility Allocation : Staff Training 05 Mar 2016 Process: 7833 Importance Of Effective Quality Management 20 Sep 2017 Process: 7845 7.1.4 Environment Of Operations 25 Sep 2017 Process: 7883 Appraisal 23 Oct 2017
ID31008	VOP 11 Equipment Control, Office, Warehouse, Pcs and Equipment Process: 5939 Responsibility Allocation : Email ISP Routing 05 Mar 2016 Process: 5941 Responsibility Allocation : Replace Main Server 07 Mar 2016 Process: 45 Responsibility Allocation : Main Server Status 16 Feb 2016 Process: 46 Responsibility Allocation : Backup Server Status 16 Feb 2016 Process: 52 Software Verification Clear Down Backup Emails 16 Feb 2016 Process: 53 Emails 16 Feb 2016 Process: 7672 Off Site Backup 09 Mar 2016 Process: 6813 Management Meeting Turnover Report 09 Mar 2016 Process: 7700 Domain Name Management 19 May 2016 Process: 7701 AWS Amazon Web Services 23 May 2016 Process: 7704 Responsibility Allocation : Computer Failure Diagnostics 24 May 2016 Process: 48 Responsibility Allocation : Internet 16 Feb 2016 Process: 49 Responsibility Allocation : Wifi 16 Feb 2016 Process: 50 Responsibility Allocation : Guest Access Wifi 16 Feb 2016 Process: 51 Responsibility Allocation : Printers 16 Feb 2016 Process: 5903 Responsibility Allocation : Weather Station 02 Mar 2016 Process: 6838 **Opera Negative Stock 10 Dec 2021 Process: 7121 Responsibility Allocation : General Computer Maintenance 09 Mar 2016 Process: 7124 Responsibility Allocation : Intrastats 09 Mar 2016 Process: 7125 Responsibility Allocation : Intrastats Urgent Problems 09 Mar 2016 Process: 7126 Intrastats Requested Page updates 09 Mar 2016 Process: 7127 Responsibility Allocation : Intrastats Unfinished in progress Processes 09 Mar 2016 Process: 7128 Responsibility Allocation : Intrastats Future Features needed 09 Mar 2016 Process: 7129 Intrastats Cross Reference Database Tables Updates 09 Mar 2016 Process: 7178 Responsibility Allocation : Systems Innovation 09 Mar 2016

	Process: 7739 Intrastats Amendment Log 12 Sep 2016 Process: 7755 Fast Hosts Invoice 08 Dec 2016 Process: 44 Secure Socket Level Certificate 16 Feb 2016 Process: 7668 Responsibility Allocation : Upgrading Intrastats ISO Quality system 09 Mar 2016 Process: 7832 Cleardown Emailed Invoices 20 Sep 2017 Process: 7823 Saftey Tester Data 02 Aug 2017
ID14696	Process: 6972 UPS Shipping Fuel Surcharge 09 Mar 2016
ID17155	VM3COP03.05 Procedures for customer returning goods on our UPS account number Process: 5879 Responsibility Allocation : Customer Returning Goods On Our UPS Account 18 Feb 2016
ID59614	Audit 15 Production Process: 7727 Audit 15 Production Viamed 24 Aug 2016 Process: 7736 Production Start Job List 03 Sep 2016 Process: 7737 Production In Production List 03 Sep 2016 Process: 7738 Production Statistics 03 Sep 2016 Process: 7775 Audit 15 Production VST 08 Feb 2017 Process: 6845 Responsibility Allocation : Quarantine Production 09 Mar 2016 Process: 6955 Production Requirements 09 Mar 2016 Process: 7169 Responsibility Allocation : Production 09 Mar 2016 Process: 7170 Responsibility Allocation : Production Production Schedule 09 Mar 2016 Process: 7171 Responsibility Allocation : Production Production Problems 09 Mar 2016 Process: 7072 Responsibility Allocation : Manufacturing Processes 09 Mar 2016
ID68045	Audit 19 Health and Safety, Working Conditions and Building Fabric Issues Process: 5941 Responsibility Allocation : Replace Main Server 07 Mar 2016 Process: 45 Responsibility Allocation : Main Server Status 16 Feb 2016 Process: 46 Responsibility Allocation : Backup Server Status 16 Feb 2016 Process: 7704 Responsibility Allocation : Computer Failure Diagnostics 24 May 2016 Process: 5856 Cleaning The Kitchen 17 Feb 2016 Process: 7729 Audit 19 Health And Saftey Viamed 24 Aug 2016 Process: 5853 Vacuuming Of The Office, Hall And Meeting Room 17 Feb 2016 Process: 5900 Cleaning Of Office Windows 25 Feb 2016 Process: 39 Enviromental Policy Document Review 16 Feb 2016 Process: 7741 Review Ethical Policy 14 Sep 2016 Process: 5878 Empty Office Bins 18 Feb 2016 Process: 5912 Responsibility Allocation : Main Recycle Bins 03 Mar 2016 Process: 7821 Controlled Waste Description And Transfer 15 Jun 2017 Process: 7820 North Yorkshire Council Waste Tranfer 15 Jun 2017 Process: 5906 Empty Paper Bins 03 Mar 2016

Process: 7805 Empty Kitchen Bins 22 May 2017
Process: 5909 Empty Warehouse Bins 03 Mar 2016
Process: 7042 Responsibility Allocation : Work Environment 09 Mar 2016
Process: 7706 Update Virus Software And Scan For Viruses 10 Jun 2016
Process: 7802 Clean Kitchen Sides 22 May 2017
Process: 7803 Dishwashing 22 May 2017
Process: 7804 Sweep Kitchen Floor 22 May 2017
Process: 7806 Watering Plants 22 May 2017
Process: 7807
Process: 7777 Audit 19 Health And Safety VST 08 Feb 2017
Process: 54 Responsibility Allocation : Gents Toilets 17 Feb 2016
Process: 5907 Hoover Warehouse 03 Mar 2016
Process: 5908 Sweep Warehouse 03 Mar 2016
Process: 5910 Clean Duckets 03 Mar 2016
Process: 5911 Clear Cardboard 03 Mar 2016
Process: 7687 Vandalism Duckets 21 Apr 2016
Process: 7698 Clean Toilets 17 May 2016
Process: 6849 First Aid 09 Mar 2016
Process: 6855 Risk Assessment HSE 09 Mar 2016
Process: 6856 Fire Alarms 09 Mar 2016
Process: 7092
Process: 56 Warehouse Outside Heating Guard 17 Feb 2016
Process: 5919 Check Out Side Drain 05 Mar 2016
Process: 5921 Clearing Water Downstairs 05 Mar 2016
Process: 7120 General Maintenance Requirements 09 Mar 2016
Process: 7742 Boiler Check 26 Sep 2016
Process: 7756 Carbon Monoxide Alarm 05 Jan 2017
Process: 48 Responsibility Allocation : Internet 16 Feb 2016
Process: 49 Responsibility Allocation : Wifi 16 Feb 2016
Process: 50 Responsibility Allocation : Guest Access Wifi 16 Feb 2016
Process: 51 Responsibility Allocation : Printers 16 Feb 2016
Process: 5903 Responsibility Allocation : Weather Station 02 Mar 2016
Process: 7121 Responsibility Allocation : General Computer Maintenance 09 Mar 2016
Process: 7178 Responsibility Allocation : Systems Innovation 09 Mar 2016
Process: 6843
Process: 7835 Electrics Need Checking 20 Sep 2017
Process: 7836 Central Heating For Winter 20 Sep 2017
Process: 7847 Health And Safety Review 26 Sep 2017
Process: 7864 ESD Work Stations 07 Oct 2017

	Process: 7867 Bandsaw Checklist 13 Oct 2017 Process: 7868 Pillar Drill Checklist 13 Oct 2017 Process: 7869 Hand Drill Checklist 13 Oct 2017 Process: 7891 Fire Alarm Evacuation Drill 25 Oct 2017 Process: 7896 Tree In Car Park 22 Dec 2017 Process: 7910 Review CCTV Warning Signs 20 Sep 2018 Process: 7928 Fire Test Points Checking 21 Feb 2019 Process: 7929 Emergency Lighting And Fire Extinguishers 21 Feb 2019 Process: 7911 Review Security Of The Special Category Personal Data 20 Sep 2018 Process: 7961 R D Room - Tidy, Empty Bins, Remove Cups. Caution Around Oxygen Supply 05 Oct 2020 Process: 7982 Check There Are No Changes To Employment Law 21 Nov 2021
ID53615	VOP 06 Measurement Control Viamed VST, Calibration, QA Stock Process: 7718 Audit 06 Calibration Viamed 24 Aug 2016 Process: 7091 Calibration Index 09 Mar 2016
ID48624	Fire risk assessment 15/17 Station Road Process: 6855 Risk Assessment HSE 09 Mar 2016
ID31032	VOP 16 Health and Safety, Company Personnel Manual Process: 7821 Controlled Waste Description And Transfer 15 Jun 2017 Process: 7820 North Yorkshire Council Waste Transfer 15 Jun 2017 Process: 6851 Review Accident Book 09 Mar 2016 Process: 7759 Health Declaration Sheet 23 Jan 2017 Process: 6849 First Aid 09 Mar 2016 Process: 6855 Risk Assessment HSE 09 Mar 2016 Process: 6856 Fire Alarms 09 Mar 2016 Process: 7092 Process: 56 Warehouse Outside Heating Guard 17 Feb 2016 Process: 5919 Check Out Side Drain 05 Mar 2016 Process: 5921 Clearing Water Downstairs 05 Mar 2016 Process: 7120 General Maintenance Requirements 09 Mar 2016 Process: 7742 Boiler Check 26 Sep 2016 Process: 7756 Carbon Monoxide Alarm 05 Jan 2017 Process: 7835 Electrics Need Checking 20 Sep 2017 Process: 7836 Central Heating For Winter 20 Sep 2017 Process: 7847 Health And Safety Review 26 Sep 2017 Process: 7867 Bandsaw Checklist 13 Oct 2017 Process: 7868 Pillar Drill Checklist 13 Oct 2017 Process: 7869 Hand Drill Checklist 13 Oct 2017 Process: 7928 Fire Test Points Checking 21 Feb 2019

ID58347	Audit 07 Handling and Storage Process: 6973 Responsibility Allocation : Stock Transfers. (QC19) 09 Mar 2016 Process: 7719 Audit 07 Handling And Storage Viamed 24 Aug 2016 Process: 7767 Audit 07 Handling And Storage VST 08 Feb 2017 Process: 5858 Opera Stock Adjustments 17 Feb 2016 Process: 5935 Stock Allocations 05 Mar 2016 Process: 6840 Process: 6850 Current Stock Levels 09 Mar 2016 Process: 6945 Missing Stock or Adjustments 09 Mar 2016 Process: 7046 Responsibility Allocation : Stock Purchasing 09 Mar 2016 Process: 7051 Responsibility Allocation : Control of nonconforming product 09 Mar 2016 Process: 7673 Check Expiry Dated Stock 09 Mar 2016 Process: 7688 Process: 7689 Move Stock From QA Shelf To Stock Shelf Monday 21 Apr 2016 Process: 7694 Move Stock From QA Shelf To Stock Shelf Tuesday 28 Apr 2016 Process: 7695 Top Up Quick Shipping Shelves 28 Apr 2016 Process: 7873 On Site Environment Review 18 Oct 2017 Process: 7866 Oxygen Cylinder Check 13 Oct 2017 Process: 7903 Empty Warehouse Depleted Sensor Bin 17 Jul 2018 Process: 7904 Check Weeee Waste Pallet And Sensor Bin 17 Jul 2018 Process: 7902 Empty Depleted Sensor Bin From The Offic 17 Jul 2018 Process: 7942 Do We Have Service Manual / QA For All Our Stock Coming In. 23 Sep 2019 Process: 7940 Review The Tom Thumb Grease Date 18 Sep 2019 Process: 7944 Sealant, Glues, Greases, Sprays, Gases And Tapes You Use In Production, Service And Repairs For Viamed And VST 09 Oct 2019
ID63048	Audit 06 Calibration Process: 7718 Audit 06 Calibration Viamed 24 Aug 2016 Process: 7766 Audit 06 Calibration VST 08 Feb 2017 Process: 7048 Control of monitoring and measuring devices 09 Mar 2016 Process: 7091 Calibration Index 09 Mar 2016
ID16995	VM3COP27.17 Complete Auto_calender Issues Process: 27 Management Reviews And Quality Audits 16 Feb 2016
ID67997	Audit 23 Analysis of Data Process: 27 Management Reviews And Quality Audits 16 Feb 2016 Process: 7733 Audit 23 Analysis Of Data Viamed 24 Aug 2016 Process: 7781 Audit 23 Analysis Of Data VST 08 Feb 2017 Process: 5877 Review Company Data 17 Feb 2016 Process: 6931 Customer Complaints 09 Mar 2016

	Process: 7839 Review VIAMED Feedback - Customer Complaints 23 Sep 2017 Process: 7838 Review VIAMED Feedback - Customer Feedback Negative 23 Sep 2017 Process: 26 Company Resources 16 Feb 2016 Process: 7070 Management Review 09 Mar 2016 Process: 7713 Review Roles And Responsibilities 17 Aug 2016 Process: 7837 Review External Parties Influencing The QMS VST / Viamed 23 Sep 2017 Process: 7840 Review VST Feedback - Customer Feedback Negative 23 Sep 2017 Process: 7841 Review VST Feedback - Customer Complaints 23 Sep 2017 Process: 7842 Review VIAMED Product Feedback Negative 23 Sep 2017 Process: 7843 Review VST Product Feedback Negative 23 Sep 2017 Process: 7071 Post Market Surveillance 09 Mar 2016 Process: 7830 Review Q.A. Failures Report 18 Sep 2017 Process: 7849 Review Product Failures New Codes 28 Sep 2017 Process: 7930 Review Flow Of Data 12 Mar 2019 Process: 7969 Weee Waste Reporting 23 Aug 2021
ID46915	VOP 10 Non Conformance, Corrective and Preventive Actions Process: 7199 Non Conformities Review Viamed 09 Mar 2016 Process: 7069 Responsibility Allocation : Corrective Actions 09 Mar 2016 Process: 7849 Review Product Failures New Codes 28 Sep 2017 Process: 7874 Review For Latest Version Med Dev 2.12. 18 Oct 2017 Process: 7264 Responsibility Allocation : VST Management Meeting Non Conformance Issues 09 Mar 2016
ID31072	VOP 08 Production, Reworks, New Production Process: 7736 Production Start Job List 03 Sep 2016 Process: 7737 Production In Production List 03 Sep 2016 Process: 7738 Production Statistics 03 Sep 2016 Process: 6845 Responsibility Allocation : Quarantine Production 09 Mar 2016 Process: 7169 Responsibility Allocation : Production 09 Mar 2016 Process: 7170 Responsibility Allocation : Production Production Schedule 09 Mar 2016 Process: 7171 Responsibility Allocation : Production Production Problems 09 Mar 2016 Process: 7072 Responsibility Allocation : Manufacturing Processes 09 Mar 2016 Process: 6962 Responsibility Allocation : VIAMED Stock Meeting Returns Overview 09 Mar 2016
ID51631	Audit 03 Design Control Process: 7716 Audit 03 Design Control Viamed 24 Aug 2016 Process: 42 Responsibility Allocation : Design Documentation 16 Feb 2016 Process: 7764 Audit 03 Design Control VST 08 Feb 2017 Process: 7043 Responsibility Allocation : Planning of product realization 09 Mar 2016 Process: 7045 Responsibility Allocation : Design and Development 09 Mar 2016 Process: 7047 Responsibility Allocation : Production and service provision 09 Mar 2016

	Process: 6942 Responsibility Allocation : Co ordination of Implementation 09 Mar 2016 Process: 7173 Responsibility Allocation : Material Generation 09 Mar 2016 Process: 5887 Review ISO/EN Documents 24 Feb 2016 Process: 7919 Send Debtors Overview To Derek 06 Dec 2018
ID63052	Audit 22 Post Market Surveillance Process: 7732 Audit 22 Post Market Surveillance Viamed 24 Aug 2016 Process: 43 Responsibility Allocation : Product Post Market Survelance 16 Feb 2016 Process: 7780 Audit 22 Post Market Surveillance VST 08 Feb 2017 Process: 6889 Responsibility Allocation : Post Market Surveilance 09 Mar 2016 Process: 7809 Pro-Active Marketing 06 Jun 2017 Process: 7810 Research Activities 06 Jun 2017 Process: 5863 Responsibility Allocation : Sales Meetings UK 17 Feb 2016 Process: 5864 Responsibility Allocation : Sales Meeting EX 17 Feb 2016 Process: 7973 VST Product Performance - Customers 27 Oct 2021 Process: 7974 VST Product Performance - Suppliers 27 Oct 2021
ID64142	Audit 11 Repairs, Servicing and Returns Process: 5898 Processing Depleted Sensors 25 Feb 2016 Process: 5879 Responsibility Allocation : Customer Returning Goods On Our UPS Account 18 Feb 2016 Process: 5857 Customer Service Logs 17 Feb 2016 Process: 7724 Audit 11 Repairs And Service Viamed 24 Aug 2016 Process: 7684 Repairs Ready For Quote 18 Apr 2016 Process: 7685 Repairs Ready For Invoice 18 Apr 2016 Process: 7690 Ship Repairs 21 Apr 2016 Process: 7748 Check Repair Orders 10 Oct 2016 Process: 7749 Check Repair Quotes 10 Oct 2016 Process: 7752 SRS Folder 22 Nov 2016 Process: 7760 Send Service Offers 31 Jan 2017 Process: 7772 Audit 11 Repairs And Service VST 08 Feb 2017 Process: 6847 Responsibility Allocation : Quarantine Repairs 09 Mar 2016 Process: 6862 Current Repairs 09 Mar 2016 Process: 7138 Non Conformance Issues Any New QC21 Forms 09 Mar 2016 Process: 7674 Check Repairs Ready For Invoice List 10 Mar 2016 Process: 7692 Responsibility Allocation : Take Complete Repair Paperwork To Office 22 Apr 2016 Process: 6916 Responsibility Allocation : Service exisiting 09 Mar 2016 Process: 6917 Responsibility Allocation : Service extension 09 Mar 2016 Process: 7823 Saftey Tester Data 02 Aug 2017 Process: 7905 Generate RMA Box, Link Items And Add Faults 17 Jul 2018 Process: 7906 Request RMA Based On The RMA Boxes 17 Jul 2018

ID75927	VOP 09 Repairs and Servicing Process: 7684 Repairs Ready For Quote 18 Apr 2016 Process: 7685 Repairs Ready For Invoice 18 Apr 2016 Process: 7690 Ship Repairs 21 Apr 2016 Process: 7752 SRS Folder 22 Nov 2016 Process: 6847 Responsibility Allocation : Quarantine Repairs 09 Mar 2016 Process: 6862 Current Repairs 09 Mar 2016 Process: 7048 Control of monitoring and measuring devices 09 Mar 2016 Process: 7674 Check Repairs Ready For Invoice List 10 Mar 2016 Process: 7814 Responsibility Allocation : Viamed Repairs 06 Jun 2017 Process: 7811 Responsibility Allocation : General Area 06 Jun 2017 Process: 7812 Responsibility Allocation : Vandagraph Repairs 06 Jun 2017 Process: 7813 Responsibility Allocation : VST Repairs 06 Jun 2017 Process: 7815 Responsibility Allocation : Product Types To Relevant Person 06 Jun 2017 Process: 7942 Do We Have Service Manual / QA For All Our Stock Coming In. 23 Sep 2019 Process: 7940 Review The Tom Thumb Grease Date 18 Sep 2019
ID75995	VOP 19 FeedBack Customer Complaints Vigilance and Notifications VST Ltd Process: 7743 Customer Complaints Paper File 26 Sep 2016 Process: 6931 Customer Complaints 09 Mar 2016 Process: 7070 Management Review 09 Mar 2016 Process: 7965 VST Feedback 29 Oct 2020
ID25632	VOP 17 Design Research and Development Process: 42 Responsibility Allocation : Design Documentation 16 Feb 2016 Process: 43 Responsibility Allocation : Product Post Market Survelance 16 Feb 2016 Process: 6975 Responsibility Allocation : Projects 09 Mar 2016 Process: 7045 Responsibility Allocation : Design and Development 09 Mar 2016
ID69314	Audit 05 Purchasing suppliers Process: 7707 Send Purchase Orders To Suppliers 13 Jun 2016 Process: 6972 UPS Shipping Fuel Surcharge 09 Mar 2016 Process: 7717 Audit 05 Purchasing Suppliers Viamed 24 Aug 2016 Process: 5850 Purchase Order Log 17 Feb 2016 Process: 7751 VST Purchase Order Log 02 Nov 2016 Process: 7765 Audit 05 Purchasing Suppliers VST 08 Feb 2017 Process: 7794 V1000 Commissions Review 30 Mar 2017 Process: 7745 UPS Invoices Viamed 06 Oct 2016 Process: 7746 UPS Invoices VST 06 Oct 2016 Process: 7747 UPS Invoices Vandagraph 06 Oct 2016 Process: 7790 Humanmed Invoice them For Previous Month 10 Mar 2017

	Process: 28 Supplier Review 16 Feb 2016 Process: 6960 Process: 5855 Purchase Order Requirements Teledyne 17 Feb 2016 Process: 5866 UPS Shipping Fuel Surcharge 17 Feb 2016 Process: 5868 Return Goods To Suppliers 17 Feb 2016 Process: 6829 Supplier Review - Outstanding orders 09 Mar 2016 Process: 6832 Supplier Review Future orders 09 Mar 2016 Process: 6848 Process: 6952 Responsibility Allocation : Lost in Shipping Claims 09 Mar 2016 Process: 6971 Responsibility Allocation : Freight Courier Cost Request 09 Mar 2016 Process: 7679 Check Stock Requirements Supplier Teledyne 18 Apr 2016 Process: 7680 Check Stock Requirements Supplier Envitec 18 Apr 2016 Process: 7681 Check Stock Requirements Supplier Posey 18 Apr 2016 Process: 7682 Check Stock Requirements Supplier Bluepoint 18 Apr 2016 Process: 7784 Check Returns Supplier Envitec 15 Feb 2017 Process: 7785 Check Returns Supplier Teledyne 15 Feb 2017 Process: 7786 Check Returns Supplier Maxtec 15 Feb 2017 Process: 7787 Check Returns All Supplier 15 Feb 2017 Process: 34 Responsibility Allocation : Insurance Is Upto Date 16 Feb 2016 Process: 7683 Check Stock For Proforma 18 Apr 2016 Process: 7882 Purchase Payments 23 Oct 2017 Process: 7956 Teledyne Stock For Vandagraph 27 May 2020 Process: 7975 Arrange Teledyne Returns 03 Nov 2021
ID75847	VOP 05 Supplier Control, Supplier Review, Purchase Orders, Supplier Returns and Rejection Process: 6972 UPS Shipping Fuel Surcharge 09 Mar 2016 Process: 28 Supplier Review 16 Feb 2016 Process: 6960 Process: 7784 Check Returns Supplier Envitec 15 Feb 2017 Process: 7785 Check Returns Supplier Teledyne 15 Feb 2017 Process: 7786 Check Returns Supplier Maxtec 15 Feb 2017 Process: 7787 Check Returns All Supplier 15 Feb 2017 Process: 7975 Arrange Teledyne Returns 03 Nov 2021
ID68263	Audit 24 Service Logs Process: 5857 Customer Service Logs 17 Feb 2016 Process: 7760 Send Service Offers 31 Jan 2017 Process: 7889 Audit 24 Servicing Viamed 24 Oct 2017
ID55437	Audit 09 Goods Inward and Product Identity Process: 5938 Responsibility Allocation : Receive Goods 05 Mar 2016

	Process: 7721 Audit 09 Goods Inward And Product Identity Viamed 24 Aug 2016 Process: 7826 Goods In Processes 06 Sep 2017 Process: 7792 Shipped Order Success Report 13 Mar 2017 Process: 7769 Audit 09 Goods Inward And Product Identity VST 08 Feb 2017 Process: 6969 Responsibility Allocation : VIAMED Stock Meeting `Goods In` Review 09 Mar 2016 Process: 57 Temporary Stock Notices 17 Feb 2016 Process: 5854 Stock FAQ Admin List 17 Feb 2016 Process: 7181 Responsibility Allocation : Product Catagories 09 Mar 2016 Process: 6894 Product Cross References 09 Mar 2016 Process: 6838 **Opera Negative Stock 10 Dec 2021 Process: 7830 Review Q.A. Failures Report 18 Sep 2017 Process: 7859 Check POR Files For Items Delivered But Not Removed From File 02 Oct 2017 Process: 7897 Daily O2 Sensors Returns 04 Jan 2018 Process: 7898 Stamp Deliveries 30 Jan 2018 Process: 7903 Empty Warehouse Depleted Sensor Bin 17 Jul 2018 Process: 7914 Proofs of Delivery 02 Oct 2018 Process: 7915 Reserve Stock Review 02 Oct 2018 Process: 7917 Human Med Purchase Order 18 Oct 2018 Process: 7923 Review Of Credits Received From Suppliers 08 Jan 2019 Process: 7943 Review Stocks Of 8000004 01 Oct 2019 Process: 7957 Warehouse Requests 29 May 2020 Process: 7962 VST Supplier QA Results 28 Oct 2020 Process: 7967 VST Stock Count For End April 01 Jul 2021 Process: 7976 Decontamination Of Incomming Products And Repairs 08 Nov 2021
ID31048	VOP 22 Picking and Packing Dispatch and Goods Out Process: 5945 Responsibility Allocation : Sending Samples 08 Mar 2016 Process: 5946 Responsibility Allocation : Sending Sale Or Returns 08 Mar 2016 Process: 7825 Responsibility Allocation : Order Picking 06 Sep 2017 Process: 5859 Review Un-shipped Parcels 17 Feb 2016 Process: 6954 Back Orders Review - By Customer 09 Mar 2016 Process: 6970 Process: 7691 Ship Sale Or Returns 21 Apr 2016 Process: 7748 Check Repair Orders 10 Oct 2016 Process: 7749 Check Repair Quotes 10 Oct 2016 Process: 7797 Check Order Are Being Picked In Priority Order 10 May 2017 Process: 6969 Responsibility Allocation : VIAMED Stock Meeting `Goods In` Review 09 Mar 2016 Process: 7860 Goods Out Picking 03 Oct 2017
ID31064	VOP 27 Software Validation Process: 52 Software Verification Clear Down Backup Emails 16 Feb 2016

	Process: 7851 **Software Validation Scan Un-QA Product To Order 23 Nov 2021 Process: 7852 Software Validation Expired Stock 01 Oct 2017 Process: 7853 Software Validation Non Sell Able Shelf 01 Oct 2017 Process: 7854 Software Validation In Production List 01 Oct 2017 Process: 7855 Software Validation - Production Lists 01 Oct 2017 Process: 7856 Software Validation Unchecked Orders 01 Oct 2017 Process: 7857 Software Validation Stock Tracking Check 01 Oct 2017 Process: 7858 Software Validation Attempt To QA Some Stock 01 Oct 2017 Process: 7861 Software Validation Of Training Documents Forced Reading 03 Oct 2017 Process: 7850 Software Validation Scan In Correct Product 01 Oct 2017 Process: 7865 Software Validation Conflicting Audits 07 Oct 2017 Process: 7870 Software Validation Non Conformance Product Risk Feedback Loop 15 Oct 2017 Process: 7879 Software Validation Scheduled Tasks And Audits 22 Oct 2017 Process: 7875 Software Validation Document Control 20 Oct 2017 Process: 7880 Software Validation Out Of Date Documents 22 Oct 2017 Process: 7881 Software Validation - Live Orders 22 Oct 2017 Process: 7892 Audit 27 Software Validation 26 Oct 2017
ID75943	VOP 20 Goods in Purchases, Returns, Repairs, Inspection / Rejection Process: 5938 Responsibility Allocation : Receive Goods 05 Mar 2016 Process: 5898 Processing Depleted Sensors 25 Feb 2016 Process: 5879 Responsibility Allocation : Customer Returning Goods On Our UPS Account 18 Feb 2016 Process: 7826 Goods In Processes 06 Sep 2017 Process: 7859 Check POR Files For Items Delivered But Not Removed From File 02 Oct 2017 Process: 7976 Decontamination Of Incomming Products And Repairs 08 Nov 2021
ID76091	Audit 14 Complaints and Corrective Actions Process: 7726 Audit 14 Complaints And Corrective Actions Viamed 24 Aug 2016 Process: 6828 Process: 7743 Customer Complaints Paper File 26 Sep 2016 Process: 7774 Audit 14 Complaints And Corrective Actions VST 08 Feb 2017 Process: 6865 Responsibility Allocation : Non Conformance Effectiveness 09 Mar 2016 Process: 7199 Non Conformities Review Viamed 09 Mar 2016 Process: 7671 Humanmed Non Conformances 09 Mar 2016 Process: 6931 Customer Complaints 09 Mar 2016 Process: 7839 Review VIAMED Feedback - Customer Complaints 23 Sep 2017 Process: 7838 Review VIAMED Feedback - Customer Feedback Negative 23 Sep 2017 Process: 7840 Review VST Feedback - Customer Feedback Negative 23 Sep 2017 Process: 7841 Review VST Feedback - Customer Complaints 23 Sep 2017 Process: 7842 Review VIAMED Product Feedback Negative 23 Sep 2017 Process: 7843 Review VST Product Feedback Negative 23 Sep 2017

	Process: 7849 Review Product Failures New Codes 28 Sep 2017 Process: 7934 Test Website Questions 02 May 2019 Process: 7965 VST Feedback 29 Oct 2020 Process: 7264 Responsibility Allocation : VST Management Meeting Non Conformance Issues 09 Mar 2016
ID77209	Process: 7728 Audit 17 Internal Audits Viamed 24 Aug 2016 Process: 7776 Audit 17 Internal Audits VST 08 Feb 2017 Process: 7972 ISO System Management Review Vst 26 Oct 2021