VOP					
Viamed Operating sub Process					
Rejected Goods – Vendor Quality Control					
Created:	27/03/06	VM3/COP/06	Issue 1		
Revised:	24 October 2017		Page 1 of 2		

1. REJECTED GOODS

- 1) Where goods fail incoming Inspection (See Procedure VM/COP/05) they will be quarantined and put on hold awaiting return to the supplier or agreed action.
- 2) If a product is rejected due to
- i) Damage in transit
- ii) Failure to meet relevant specification
- iii) Failure to meet IEC601 safety specification.

It is placed in "Reject Return To Supplier" area suitably labelled

- 3) Where appropriate, a 'New Return Shipment' will be raised and Emailed to the supplier as pre-notice of a pending return, request for a returns number, collection, or sent with the goods.
- 4) The Return Shipments are uniquely numbered and is logged in intrastats.

2. .<u>VENDOR QUALITY CONTROL</u>

1) Responsibilities

The Managing Director will be responsible for ascertaining whether Suppliers/Subcontractors performance is satisfactory or unsatisfactory based on the following:-

For each Product

- (1) Rejects as a result of non-conformance to purchase order.
- (2) Rejects as a result of damage in transit.
- (3) Incorrect or lack of correct documentation.
- (4) Incorrect quantities delivered.
- (5) Customer complaints.
- (6) Length of time taken in correcting situation
- (7) Corrective action will be taken by the Managing Director

1) Quality Record

Each Supplier/Sub Contractors quality records will be kept in the Suppliers File and records kept of any action taken against them

2) Concessions

All concessions on quality will be at the Managing Directors sole discretion and only if the product still complies with the minimum acceptable quality required.

3) Non Conformity Review

Non conformity of any product must be immediately reviewed by the managing director and QA personnel before release

4) Corrective Action

Where possible existing procedures: Repairs VOP09 and Customer Complaints VOP19 should be used

5) Preventetive Action

Manufacturers and suppliers should be informed by Fax or email of any anomalies in product or installations found in routine service or maintenance. If potentially serious a customer complaint should be raised.

	VOP				
	Viamed	Operating sub	Process		
Rejected Goods – Vendor Quality Control					
Created:	27/03/06	VM3/COP/06	Issue 1		
Revised:	24 October		Page 2 of 2		
	2017				