

<b>VOP</b>			
<b>Operating sub Process</b>			
<b><u>SUPPLIER CONTROL</u></b>			
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<b><u>QC06</u></b>			

### **SCOPE**

This procedure is established to describe the system used within the company for the control of purchasing of product and evaluation / assessment of suppliers and sub-contractors. It is used in conjunction with the individual sub procedures, which show the relevant information necessary.

### **RESPONSIBILITIES**

It is the responsibility of the Managing Director, to ensure that the contents of this procedure, and related procedures, are adhered to. It is the responsibility of the relative supervisors to ensure that their requirements are made clear. The Warehouse Controller is responsible for the ongoing evaluation of suppliers / sub-contractors.

### **PURCHASING**

A warehouse request is submitted by a member of staff. This is picked up by the Warehouse controller, they then generated a Purchase order POR in our accounts package. To cover the supply of products for warehouse stock, or immediate customer delivery.

The purchase order is raised for goods or materials as required, and the details entered into the correct supplier on the accounts package. The printed purchase order form is then forwarded to a Director for approval. Once it has been approved it can then be sent to the supplier by the office.

Requisitions for Orders may be initiated by:

Engineers – stock needed for production or repairs.

Office staff – when they check stock for customers order and it show we will not have sufficient stock.

Warehouse controller – to bring stock items up to minimum level or against goods not already in stock or already ordered, to supply customer orders in hand.

All POR are countersigned by a Director or produced by a director.

Purchase orders are processed by the following means;

- Addressee abroad – By Fax/Email
- Non urgent orders By – Post/Email
- Urgent orders – By Fax/Email
- Extremely urgent orders – By Telephone/Email, (the POR still needs to be printed and signed off)

Purchase Order Number is generated automatically. Purchase Orders contain all relevant data to the supplier, part numbers or product codes, description and quantity.

A printed copy of the POR must be made for company records and supplier confirmation. The Purchase order POR is filed in the goods in department in the POR

file. So that when the goods come in, they can be checked against our original paperwork.

The Purchase Order must specify any requirements for inspection and tests, and any certification required, include special instructions for packaging, labelling or delivery. Additional information where appropriate is advised e.g.

- Quoted price when applicable.
- “Urgent Please” if supply is required urgently.
- “Partial shipment acceptable/not acceptable”.
- Preferred method of shipment.
- If supplier needs to know for traceability of specified instruments or service. We can give the customer or instrument details.
- Requirement on POR for serial numbers/ LOT numbers / batch numbers for traceability.
- Conformity Certification needs to be requested from the supplier when required.

Purchase Orders will be reviewed and authorised by the Managing Director, Director or Warehouse Controller.

It is the responsibility of the individual raising the purchase order to ensure that all the relevant and correct information is entered onto the purchase order. It is the responsibility of the person receiving the Intrastats task to ensure the copy purchase orders are filed accordingly.

### **ASSESSMENT OF SUPPLIERS / SUB-CONTRACTORS**

All products provided to us under cover of its ISO 9001/13485 must be procured from a Quality Assured Source, or is internally QA / Verified. When a customer orders products, in accordance with the companies Approved Supplier List, then a quality assured source will be:-

- Companies listed in the DTI Quality Assurance Register of assessed companies current issue, and who supply within their declared scope.
- Companies assessed by certification bodies outside the UK to equivalent systems.
- A Company specifically nominated by a customer in his order will also be considered a Quality Assured Source for that particular customer, or where the customer has specified the product by brand or specific part number.

### **APPROVAL OF SUPPLIERS**

New products will be considered by the Managing Director, by assessment of suppliers capability to meet the required quality standard. At an acceptable delivery date, to verify the firm maintains batch segregation, inspection and test records. A Supplier/Sub contractor questionnaire (Form QC06) will be used to approve suppliers.

A Register of Approved Suppliers/Sub contractors will be maintained by Managing Director or his chosen representative in Intrastats, which will distinguish between Quality Assured and non-Quality Assured Sources.

In certain instances, commercial considerations may result in the use of a supplier not included on the approved supplier list. In these instances the level of goods inwards inspection is to a higher standard until such time as the supplier has passed assessment and been entered onto the approved register.

At least once per year all suppliers will be re validated or re-assessed.

A register of current suppliers is available and maintained in Intrastats. This register is monitored and evaluated on a regular basis, and formally reviewed annually for the management review. The determination of supplier suitability is made on the basis of Third party approval, historical acceptable performances, pricing, delivery or other such factors as may be deemed appropriate.

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Information received from all departments, on an annual basis, will be taken into consideration for review of this register.

#### **CERTIFICATION**

Conformity/Test Certificates need to be obtained from the supplier when they are offered by us, to the customer. Information is copied from manufacturer/supplier certification. The original certificate is sent to the customer and a copy retained with the customers ORD paperwork.

#### **PRODUCTS OUTSIDE THE REGISTRATION**

Any product which is not included in the Registration will be clearly identified in the covering documentation so that there can be no misunderstanding on the part of the customer.

Similarly if a supplier has not been approved as specified Approval of Suppliers, their products must not be sold without them being clearly identified and confirmed in the covering documentation, as follows - "This item procured from a non ISO 9001 source".