

Supply Chain New Product Introduction (SC02)

Date	Version	
27/04/17	2	
Purpose		
To identify the key activities in managing products throughout its lifecycle, including; <ol style="list-style-type: none"> 1. New Product Introduction 2. Product Obsolescence 		
Who should read this document?		
All those that are involved in introducing, using or changing products should read this document.		
Key messages		
SOPs aim to achieve efficiency, quality output and uniformity of performance, while reducing miscommunication and failure to comply to industry or Trust regulations.		
Accountabilities		
Production	Supply Chain Manager	
Review and approval	Scan4Safety GTIN/P2P Project Board	
Ratification	Chief Procurement Officer	
Dissemination	Head of Supply Chain & E-Procurement	
Compliance	Records & Information Forum	
Links to other policies and procedures		
Supply Chain Inventory Management Policy (SC010v.1)		
Version History		
1	21 st March 2017	Ratified by Chief Procurement Officer and published Trust-wide
2	2 nd May 2017	Ratified by Chief Procurement Officer and published Trust-wide
Last Approval		Due for Review
May 2017		May 2020

The Trust is committed to creating a fully inclusive and accessible service. By making equality and diversity an integral part of the business, it will enable us to enhance the services we deliver and better meet the needs of patients and staff. We will treat people with dignity and respect, promote equality and diversity and eliminate all forms of discrimination, regardless of (but not limited to) age, disability, gender reassignment, race, religion or belief, sex, sexual orientation, marriage/civil partnership and pregnancy/maternity.

An electronic version of this document is available on the Trust Documents Network Share Folder (G:\TrustDocuments). Larger text, Braille and Audio versions can be made available upon request.

Standard Operating Procedures are designed to promote consistency in delivery, to the required quality standards, across the Trust. They should be regarded as a key element of the training provision for staff to help them to deliver their roles and responsibilities.

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Standard Operating Procedure (SOP) Supply Chain Product Lifecycle

1 Purpose and Scope

1.1 Introduction

The purpose of this SOP is to identify the key activities involved in the introduction, change and removal of a product within the Trust.

This procedure is applicable to all those involved in the ordering, using and managing of product inventory within the Trust.

It should be applied across the Trust, with particular relevance to Procurement and Supply Chain functions.

1.2 Definitions

- **SOP** – Standard Operating Procedure
- **SCM** – Supply Chain Manager
- **CPO** – Chief Procurement Officer
- **DoF** – Director of Finance
- **HSCEP** – Head of Supply Chain & E-Procurement
- **SCTL** – Supply Chain Team Leader
- **SCT** – Supply Chain Team
- **RCA** – Root Cause Analysis
- **SSCM** – Senior Supply Chain Management
- **NPI** – New Product Introduction
- **PO** – Purchase Order
- **IMS** – Inventory Management System

1.3 Regulatory background

n/a

1.4 Key Duties

Key functional duties are outlined on process flow chart in section '2. Procedure to follow'.

1.5 Monitoring and assurance

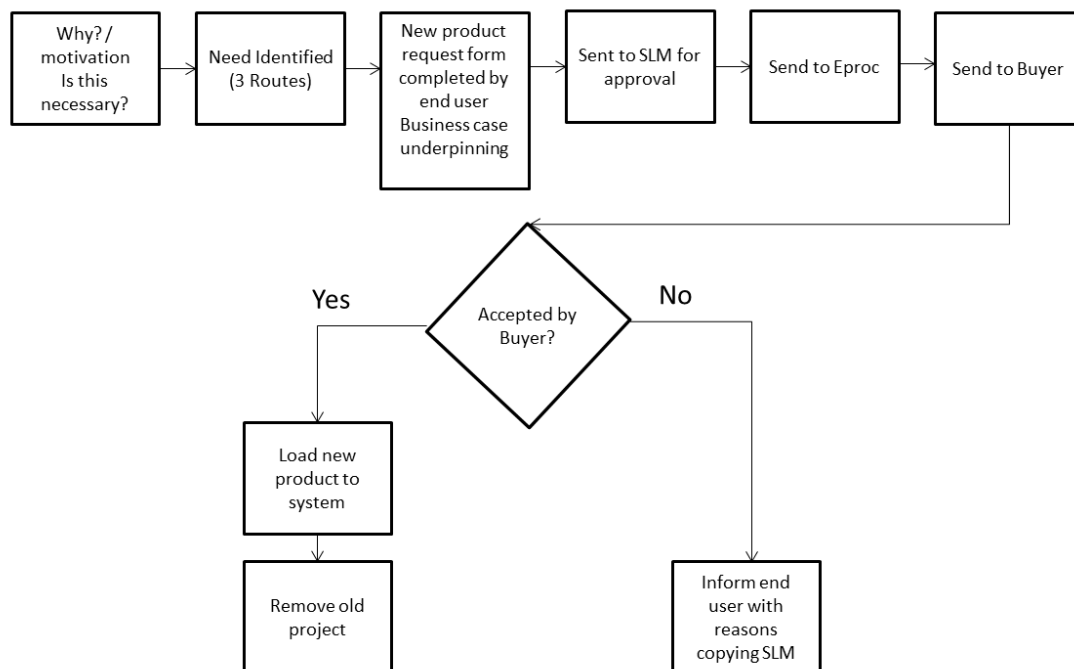
- Accountability for adherence to this procedure will be monitored by the Senior Supply Chain Management (SSCM).
- Each Category Manager or Buyer will be responsible for ensuring this procedure is followed operationally
- Adherence to the identified process flow will be monitored through monthly analysis of new part numbers and identification of the part number it replaces (if applicable)

- Supply Chain Team perform Root cause analysis (RCA) for the following;
 - New parts which have not followed this procedure
 - Obsolete parts which have not followed this procedure
- For every RCA conducted, findings to be reported to the Senior Supply Chain Management
- Supply Chain Team have a responsibility to support the implementation of change

2 Procedure to Follow

2.1 New Product Introduction (NPI)

Current New Product Request Process



The above process should be owned by the relevant Buyer from Procurement. Supply Chain and Customer Representatives will have specific responsibilities throughout the process (see “How to request a new product” notification in appendix) and success will depend on these tasks being delivered and managed in a coordinated and timely manner. The Buyer will develop and own the integrated project plan for managing NPI and Obsolescence activity.

For new clinical devices reference TRW.HGV.POL.229.5 The introduction of new clinical devices and procedures policy which includes “The introduction of new clinical devices and procedures policy flow chart”

3 Document Ratification Process

3.1 The design and process of review and revision of this procedural document will comply with The Development and Management of Trust Wide Documents.

The review period for this document is set as default of three years from the date it was last ratified, or earlier if developments within or external to the Trust indicate the need for a significant revision to the procedures described.

This document will be approved and ratified by the CPO.

Non-significant amendments to this document may be made, under delegated authority from the CPO, by the nominated author.

Significant reviews and revisions to this document will include a consultation with named groups, or grades across the Trust. For non-significant amendments, informal consultation will be restricted to named groups, or grades who are directly affected by the proposed changes

3.2 Dissemination and Implementation

Following approval and ratification, this procedural document will be published in the Trust's formal documents library and all staff will be notified through the Trust's normal notification process, currently the 'Vital Signs' electronic newsletter.

Document control arrangements will be in accordance with The Development and Management of Trust Wide Documents.

The document author(s) will be responsible for agreeing the training requirements associated with the newly ratified document with the CPO and for working with the Trust's training function, if required, to arrange for the required training to be delivered.

4 Reference Material

n/a

How to Request a New Product

Please follow the below steps:

- **Brand new item...**

Please complete and submit a 'Eproc New Product Request Form' to your Service Line Management Team. NPRF available via Staffnet link shown below.

<http://staffnet.plymouth.nhs.uk/Departments/OtherSupportServices/Finance,ProcurementandPerformanceInformation/Procurement/Systems.aspx>

- **Add a variant of an existing store room stock...**

Please contact your store person who shall complete and submit a 'Internal Eproc New Product Request Form' to the Procurement Team.

- **Procurement driven change...**

The Procurement team shall complete and submit a 'Internal Eproc New Product Request Form' to eprochelpdesk.

- **Supplier driven change...**

Please contact the Procurement Team. (Example: Item no longer available)

Area: Supply Chain Team Communication Board

Responsible Team: Supply Chain Team

1st Contact: Dave Knight (Ext: 32720)

2nd Contact: Paul Cady (Ext: 30098)



Plymouth Hospitals
NHS Trust

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