

COGPAC8

Control packing operations for pilots and clinical trials



Overview

This unit describes the competences required to control a packing operation for a pilot or clinical trial.

To perform competently, you will need to show that you can operate in a range of conditions. You will need to demonstrate therefore that you can deal effectively with the following:

- 1 checks of the packs (primary, secondary or tertiary), product security and safety
- 2 procedures relating to legal requirements and quality standards
- 3 problems associated with documentation, services, area and equipment and faults in production quality
- 4 documentation or packing instructions, written packing records and computer-based packing records
- 5 handover procedures

During this work you must take account of the relevant worksite operational requirements, procedures and safe working practices **AS THEY APPLY TO YOU**.

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Performance criteria

You must be able to:

- P1 use the appropriate personal protective equipment (PPE) and/or product-specific equipment/clothing
- P2 pack the correct quantity of products to the correct specification
- P3 segregate and label correctly materials and packs which do not meet the specification
- P4 maintain the required output rates
- P5 keep the work area clean, tidy and secure in compliance with procedures
- P6 prepare the packs correctly for transfer
- P7 identify and deal with problems according to the procedure
- P8 complete the required documentation accurately and legibly
- P9 maintain the correct operational status when you have taken over responsibility
- P10 report the current operational status accurately within the acceptable time limits, at the correct time and place
- P11 follow the correct procedures when handing over to and receiving from someone else

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Knowledge and understanding

You need to know and understand:

- K1 which personnel protective equipment (PPE) is appropriate and why
- K2 any specialised protective equipment or clothing needed for the product and its use
- K3 the critical factors affecting product quality (e.g. temperature, humidity etc.)
- K4 any specific product handling requirements and sources of information (e.g. hazard sheets)
- K5 how to interpret and identify packing instructions
- K6 what are the expected levels of cleanliness for the work area
- K7 what is the procedure for dealing with spillages
- K8 how to monitor product security/integrity during packing
- K9 the procedure for segregating materials and packs
- K10 what are the labelling procedures
- K11 the rate at which you are expected to work
- K12 how to monitor and maintain the levels of materials
- K13 how to make adjustments according to the procedure
- K14 how to prepare packs for transfer
- K15 how to handle and dispose of non-recoverable packs and materials
- K16 the procedures for reporting problems
- K17 the methods of dealing with problems
- K18 what information is to be transferred and in what format
- K19 the requirements of the handover procedure (both giving and receiving)
- K20 how to maintain product security/integrity during the handover
- K21 how to identify and emphasise significant information
- K22 the operational quality standards required

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