

Australian Government

# FBPPHM3002 Operate a pharmaceutical production process

Release: 2

## FBPPHM3002 Operate a pharmaceutical production process

Release	Comments	
Release 2	This version released with FBP Food, Beverage and Pharmaceutical Training Package Version 4.1.	
Release 1	This version released with FBP Food, Beverage and Pharmaceutical Training Package Version 2.0.	

#### **Modification History**

# Application

This unit of competency describes the skills and knowledge required to setup, operate monitor, adjust and shut down a production process in a pharmaceutical manufacturing facility.

The unit applies to individuals who apply operating principles to the production process. Individuals work under broad direction and take responsibility for their own work.

No licensing, legislative or certification requirements apply to this unit at the time of publication.

## Pre-requisite Unit

Nil

## **Unit Sector**

Pharmaceutical (PHM)

Elements	Performance Criteria
Elements describe the essential outcomes.	Performance criteria describe the performance needed to demonstrate achievement of the element.
1. Receipt materials and components	<ul><li>1.1 Confirm incoming goods correspond to workplace documentation</li><li>1.2 Clean and label containers with prescribed data, according to workplace procedures</li></ul>
	1.3 Quarantine incoming goods including release and reject status according to Good Manufacturing Practice (GMP) requirements and

#### **Elements and Performance Criteria**

Elements	Performance Criteria		
Elements describe the essential outcomes.	Performance criteria describe the performance needed to demonstrate achievement of the element.		
	workplace procedures		
	1.4 Identify and report deviations, unusual events and non-conformances according to GMP and workplace procedures		
2. Set up the production	2.1 Confirm equipment and materials meet production requirements		
process for operation	2.2 Confirm cleaning requirements and equipment status		
	2.3 Select and fit personal protective equipment and contamination prevention clothing according to workplace procedures		
	2.4 Enter processing and operating parameters according to safety and production requirements		
	2.5 Check and adjust equipment performance		
	2.6 Conduct pre-start checks according to workplace procedures		
3. Dispense materials	3.1 Deliver materials in required quantities and sequence according to batch and production requirements		
	3.2 Record dispensed material, including weight or volume according to batch and production requirements		
	3.3 Label dispensed materials for each batch and stage according to production requirements		
4. Operate and monitor the production process	4.1 Start up, monitor and control production process to maintain process within required limits		
	4.2 Identify and report out of limit products or processes according to workplace procedures		
	4.3 Maintain work area according to workplace cleaning standards		
	4.4 Conduct production process according to safety and environmental requirements		
	4.5 Complete documentation according to workplace procedures		
5. Hand over the	5.1 Perform handover according to workplace procedures		
production process	5.2 Inform handover production team of process and related equipment status at completion of handover		
6. Shut down the process	6.1 Confirm the workplace procedures for shutting down the process		
	6.2 Complete end-of-batch procedures according to batch instructions and workplace procedures		
	6.3 Safely shut down the process		

Elements	Performance Criteria	
Elements describe the essential outcomes.	Performance criteria describe the performance needed to demonstrate achievement of the element.	
	6.4 Complete records according to workplace procedures	

# **Foundation Skills**

This section describes those language, literacy, numeracy and employment skills that are essential for performance in this unit of competency but are not explicit in the performance criteria.

Skill	Description	
Reading	• Identify relevant information from workplace documentation and interpret requirements for the pharmaceutical production process	
Writing	Complete workplace documentation using appropriate language and in required format	
Numeracy	Interpret material and product specifications	

# **Range of Conditions**

This section specifies different work environments and conditions that may affect performance. Essential operating conditions that may be present (depending on the work situation, needs of the candidate, accessibility of the item, and local industry and regional contexts) are included.

Cleaning requirements must include:	<ul> <li>area or line clearance</li> <li>full or partial clean</li> <li>automated or semi-automated or manual cleaning of equipment</li> <li>sanitation or sterilisation.</li> </ul>	
Equipment status must include:	<ul> <li>calibrated</li> <li>clean</li> <li>clean/dirty hold time</li> <li>in use</li> <li>ready to use.</li> </ul>	
Pre-start checks must include:	carrying out required area or line clearances inspecting equipment condition to identify signs of wear confirming all safety equipment is in place and operational	

	<ul> <li>confirming that equipment is clean or sanitised</li> <li>confirming that equipment is correctly configured for proce requirements</li> </ul>	
Items to monitor must include:	•	environment product appearance volume or weight.

# **Unit Mapping Information**

Code and title current version	Code and title previous version	Comments	Equivalence status
FBPPHM3002 Operate a pharmaceutical production process Release 2	FBPPHM3002 Operate a pharmaceutical production process Release 1	Minor updates to Range of Conditions for clarity Foundation skills refined	Equivalent

### Links

Companion Volume Implementation Guides are found in VETNet: https://vetnet.gov.au/Pages/TrainingDocs.aspx?q=78b15323-cd38-483e-aad7-1159b570a5c4