# **Modification history**

Release	Comments
Release 2	This version released with FBP Food, Beverage and Pharmaceutical Training Package Version 3.0.
Release 1	This version released with FBP Food, Beverage and Pharmaceutical Training Package Version 2.0.

FBPPHM4003	Facilitate contamination control
Application	This unit of competency describes the skills and knowledge required to identify and implement contamination control measures in a pharmaceutical manufacturing facility.
	The unit applies to individuals with specialised skills and knowledge of Good Manufacturing Practice (GMP) requirements who are responsible for overseeing the implementation of cleaning, sanitation, change control and validation programs within pharmaceutical manufacturing operations and have responsibility for the output of others. This includes applying and communicating non-routine technical solutions to predictable and unpredictable problems.
	No occupational licensing, legislative or certification requirements apply to this unit at the time of publication. Error! Use the Home tab to apply AFSA AR Code to the text that you want to appear here.
Prerequisite 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.
Review contamination risks and related control	1.1 Identify hazards that could present contamination risks by type, origin and product association
measures to meet GMP requirements	1.2 Review hazards and risk control measures to confirm they meet GMP requirements
·	Conduct risk assessment according to GMP requirements and workplace procedures     Confirm control measures
Implement procedures used to control risk of cross contamination	2.1 Identify effective barriers and control systems to minimise risk of cross-contamination according to GMP requirements and workplace procedures 2.2 Conduct in-process and environmental monitoring according to GMP requirements and workplace procedures 2.3 Validate cleaning processes according to workplace procedures 2.4 Confirm line clearance procedures 2.5 Ensure personal hygiene and conduct of personnel in work area meets GMP requirements
	2.6 Ensure operators have the skills and knowledge required to apply contamination control measures

Foundation	Skills
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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.

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Skill	Description	
Reading	<ul> <li>Interpret key information from in-process and environmental monitoring test results</li> <li>Interpret key information about GMP requirements in workplace procedures relevant to facilitating contamination control</li> </ul>	
Writing	Record contamination information in document management systems	
Oral communication	Use appropriate language and communication skills to ensure that contamination management procedures are understood and implemented	
Numeracy	Monitor data to determine instances of contamination according to GMP requirements	

Unit Mapping Information			
Code and title current version	Code and title previous version	Comments	Equivalence status
FBPPHM4003 Facilitate contamination control Release 2	FBPPHM4003 Facilitate contamination control Release 1	Foundation skills amended	Equivalent unit

Links	Companion Volumes, including Implementation Guides, are available at
	VETNet https://vetnet.gov.au/Pages/TrainingDocs.aspx?q=78b15323-cd38-
	483e-aad7-1159b570a5c4

TITLE	Assessment requirements for Error! Use the Home tab to apply
	AFSA Unit Code to the text that you want to appear here. Facilitate
	contamination control

#### **Performance Evidence**

An individual demonstrating competency in this unit must satisfy all the elements and performance criteria of this unit.

There must be evidence that the individual has facilitated contamination control in at least one workplace, including:

- identified and investigated occurrences that could present contamination hazards and risks by:
  - type
  - origin
  - product association
- · confirmed contamination control measures, including:
  - critical limits
  - · monitoring and recording requirements
- interpreted Good Manufacturing Practice (GMP) requirements and workplace procedures related to contamination control
- used document management systems to:
  - access and review documents regarding cross-contamination
  - monitor and record data according to GMP requirements
  - ensure procedures are understood and implemented
- reviewed workplace documents to determine contamination control measures, including:
  - equipment drawings
  - piping and instrumentation diagrams
  - · process flow charting
- monitored gowning, cleaning, access and refresher training according to workplace procedures
- provided training and support to others in work area to implement contamination control according to GMP requirements and workplace procedures.

### **Knowledge Evidence**

An individual must be able to demonstrate the knowledge required to perform the tasks outlined in the elements and performance criteria of this unit. This includes knowledge of:

- types of contamination, including:
  - microbiological
  - chemical
  - physical
- microbiological limits, monitoring methods and reporting and recording formats and requirements
- line clearance procedures, roles and responsibilities
- personal hygiene and clothing requirements, including:
  - informing team leader or supervisor of reportable illness
  - · removal of jewellery
  - removal of makeup
  - personal clothing use, storage and disposal
  - personal clothing and footwear requirements for working in and moving between work areas
  - decontamination
  - laundering
- GMP requirements and workplace procedures relevant to contamination control
- environmental issues and workplace controls relevant to contamination control, including:
  - the principles of workflow design to minimise risk of contamination
  - ventilation system requirements
  - · production facility segregation requirements
  - storage requirements of raw materials prior to their use in manufacture
  - · time limits and conditions for storing finished products prior to packaging
  - · requirements relevant to product range
- workplace cleaning standards and responsibilities, including:

## **Knowledge Evidence**

- types of cleaning agents and what they do
- waste collection
- recycling, safe handling and disposal of different types of waste
- · safe handling and disposal of hazardous waste
- requirements for completion of workplace documentation.

#### **Assessment Conditions**

Assessment of skills must take place under the following conditions:

- physical conditions:
  - a pharmaceutical manufacturing workplace or an environment that accurately represents workplace conditions
- · resources, equipment and materials:
  - document management system
  - · equipment drawings, piping and instrumentation diagrams and process flow charts
- specifications:
  - · recording requirements and procedures
  - workplace procedures related to contamination control
  - · GMP requirements related to contamination control
  - workplace cleaning procedures
- relationships:
  - · team members.

Assessors of this unit must satisfy the requirements for assessors in applicable vocational education and training legislation, frameworks and/or standards.

Links	Companion Volumes, including Implementation Guides, are available at VETNet
	https://vetnet.gov.au/Pages/TrainingDocs.aspx?q=78b15323-cd38-483e-aad7-
	1159b570a5c4