Modification history

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| Release | Comments |
| Release 1 | This version released with FBP Food, Beverage and Pharmaceuticals Training Package Version 2.0. |

| FBPPHM4003 | Facilitate contamination control |
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| 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. |
| Prerequisite Unit | Nil |
| Unit Sector | Pharmaceutical (PHM) |

| Elements | Performance Criteria |
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| Elements describe the essential outcomes. | Performance criteria describe the performance needed to demonstrate achievement of the element. |
| 1. Review contamination risks and related control measures to meet GMP requirements | 1.1 Identify hazards that could present contamination risks by type, origin and product association1.2 Review hazards and risk control measures to confirm they meet GMP requirements1.3 Conduct risk assessment according to GMP requirements and workplace procedures1.4 Confirm control measures |
| 2. 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 procedures2.2 Conduct in-process and environmental monitoring according to GMP requirements and workplace procedures2.3 Validate cleaning processes according to workplace procedures2.4 Confirm line clearance procedures2.5 Ensure personal hygiene and conduct of personnel in work area meets GMP requirements2.6 Ensure operators have the skills and knowledge required to apply contamination control measures |

| Foundation SkillsThis 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 | * Interpret key information from in-process and environmental monitoring test results
* Interpret key information in GMP requirements and workplace procedures relevant to facilitating contamination control
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| Writing | * Record contamination information in document management systems
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| Numeracy | * Monitor data to determine instances of contamination according to GMP requirements
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| Navigate the world of work | * Apply electronic document management systems
* Monitor personnel adherence to GMP requirements and workplace procedures
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| Interact with others | * Use appropriate language and communication skills to ensure that contamination management procedures are understood and implemented within the work area
* Report GMP requirements for contamination management to relevant personnel using required communication method
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| Unit Mapping Information |
| Code and title current version | Code and title previous version | Comments | Equivalence status |
| FBPPHM4003 Facilitate contamination control | FDFPH4003A Facilitate contamination control | Updated to meet Standards for Training Packages. | Equivalent unit |

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

| TITLE | Assessment requirements for Facilitate contamination control |
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| 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, on at least one occasion, the individual has facilitated contamination control, 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 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 (P&IDs)
* 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.
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| Knowledge Evidence |
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| 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:* understand the 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:
* making team leader or supervisor aware 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:
* 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.
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| Assessment Conditions |
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| 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 (PI&Ds) and process flow charting
* 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. |

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