

Product Preparation PLAR Requirement - Sterile Compounding

**Verification of Competency in Sterile Product Preparation**

**Name of PLAR Candidate (please print):**

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**Pharmacy Regulatory ID Number:**

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**Delivery institution where PLAR Challenge Exam will be written:**

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**Assignment:** Participants must demonstrate safe and accurate preparation of a sterile parenteral admixture in a Laminar Air Flow Hood (LAFH) with emphasis on aseptic technique and meeting the standards established by the Canadian Society of Hospital Pharmacists (CSHP), USP Chapter 797 and Good Manufacturing Practice (GMP)

Competency	Competency Demonstrated	Competency Not Demonstrated	Student's Initials	Assessor's Initials
<b>Set Up</b>				
Selected appropriate ingredients				
Selected appropriate equipment				
Accurately calculated quantities				
Prepared labels prior to entering the LAFH				
Positioned supplies appropriately in the hood				
<b>Operator Preparation</b>				
Demonstrated appropriate hand washing technique				
Selected and put on appropriate personal protective equipment (gloves, gown, mask, head cover, etc.)				
<b>Aseptic Technique</b>				
Performed appropriate disinfection procedures – e.g. gloves, puncture sites				
Performed procedures 6 inches within the LAFH and 3 inches above the work surface				
Ensured critical sites were in direct airflow				
Demonstrated appropriate needle safety techniques				
Placed needle caps on alcohol swabs during manipulations				
Re-sprayed or wiped gloves when re-entering the hood (after leaving to retrieve additional supplies)				

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Wiped spills immediately with either an alcohol swab or with a gauze soaked with 70% alcohol				
<b>Specific Procedures to be Demonstrated</b>				
<b>Withdrawal from a Multi-Use Vial:</b> Demonstrated appropriate technique				
<b>Withdrawal of a Solution from an Ampoule:</b> Demonstrated appropriate technique				
<b>Injection of a Solution into an I.V. bag:</b> Demonstrated appropriate technique				
<b>Reconstitution of a Powder:</b> Demonstrated appropriate use of technique with a non-vented vial				
Demonstrated appropriate use of technique with a vented needle				
<b>Final Steps</b>				
Inspected the finished product before performing the verification				
Followed protocols for labeling and sealing product				
Verified appropriateness of product to prescription				
Completed required documentation				
Cleaned preparation area and equipment appropriately				
Removed and discarded personal protective equipment appropriately				

**Assessment completed by:**

Assessor's Signature: \_\_\_\_\_

Please Print Name and Title: \_\_\_\_\_

Business Phone Number: \_\_\_\_\_

Pharmacy Regulatory Authority I.D. Number: \_\_\_\_\_

Date of completion of assessment: \_\_\_\_\_

Name/Address of Pharmacy (or other site) where assessment was completed:  
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