



DATE:

I. ABOUT YOUR COMPANY

**Fields required to be filled out*

1. NAME*

2. DESIGNATION*

3. COMPANY*

4. ADDRESS*

5. EMAIL*

6. WEBSITE

7. PHONE NUMBER AND EXTENSION*

8. FAX

9. YOU WORK FOR*

(Please Tick)

- End User/Facility Owner
- Cleanroom Builder/Contractor
- Lab Builder/Contractor
- Distributor

10. EXISTING ESCO EQUIPMENT*

11. REPEATED ORDER*

- Yes, SN: No

II. PROJECT INFORMATION

**Fields required to be filled out*

12. URS Available*

- Yes (please attach document)
 No

13. Industry*

- Hospital Pharmacy
- Pharmaceutical
- Outsourcing Facility
- Chemicals
- Research and Development
- Others, please specify:

14. Name of Project

15. Project Location
(City, Country)*

16. Deadline of submission for Tender*

17. Timeline of Purchase

18. Timeline of Installation*

19. Application/s*

- Sterile Pharmacy Compounding
- Non-sterile Pharmacy Compounding
- Hazardous Drug Compounding
- Radiopharmaceutical Compounding

For radiopharmaceutical applications, please refer to and use the "Radiopharmacy Equipment Questionnaire".

- Sterility Testing
- Research and Development
- Others:

20. Type of Secondary Engineering Control* (Room where to place the cabinet)

- Cleanroom
 - ISO Class 7
 - ISO Class 8
 - Others, please specify:

- Segregated Compounding Area
- Others, please specify:
- Room Dimension (W x D x H)
 - Height Clearance:
 - Door Dimension and Clearance :

21. Provide Site Plan/Floor Layout so that Esco can verify clearances are Sufficient for Installation/ Maintenance Access

Please attach the site plan/floor layout together with this questionnaire.

22. What standards do you follow?

23. For Sterile Pharmacy Compounding

- Type of Preparation
 - Total Parenteral Nutrition
 - Antibiotic Compounding
 - Others:

	<ul style="list-style-type: none"> • Equipment Needed for the Process: <input type="text"/> • Will you carry out sterility testing using a sterility test pump? <input type="checkbox"/> Yes, brand/model: <input type="text"/> <input type="checkbox"/> No
<p>24. For Non-sterile Pharmacy Compounding</p>	<ul style="list-style-type: none"> • Type of Preparation <input type="checkbox"/> Tablets/Capsules <input type="checkbox"/> Oral Solutions/Suspensions <input type="checkbox"/> Dermatological Preparations <input type="checkbox"/> Others: <input type="text"/> • Equipment Needed for the Process: <input type="text"/>
<p>25. For Hazardous Drug Compounding</p>	<ul style="list-style-type: none"> • Type of Hazardous Drug: <input type="checkbox"/> Non-sterile HD <input type="checkbox"/> Sterile HD <input type="checkbox"/> Both • Do you handle these drugs or any other volatile drugs? <i>(Carmustine, Cyclophosphamide, Doxorubicin, Ifosfamide, Meclorothamine, Thiotepa)</i> <input type="checkbox"/> Yes <input type="checkbox"/> No
<p>26. Protection (Tick All That Apply)</p>	<ul style="list-style-type: none"> <input type="checkbox"/> Operator protection <input type="checkbox"/> Product protection <input type="checkbox"/> Environmental protection
<p>27. Level of Need</p>	<ul style="list-style-type: none"> <input type="checkbox"/> Have an approved budget (indicate: <input type="text"/>) <input type="checkbox"/> Preparing to submit a budget for approval <input type="checkbox"/> Gathering information for future reference

III. CABINET SPECIFICATIONS INFORMATION

*Fields required to be filled out

28. Internal Width*

29. Internal Depth*

30. Internal Height*

31. Type of Cabinet*

- Open Front
 Closed-system/Isolator

32. Pressure Mode*

- Positive Pressure
 Negative Pressure
 Required Pressure, please specify per chamber:

33. Airflow Circulation*

- Recirculating
 Total Exhaust

34. Airflow Pattern*

- Unidirectional
 Turbulent

35. Construction Material

(Indicate if interior or exterior material of construction)

Specify the preferred material of construction:

Internal:

External:

- Antimicrobial Powder-Coated ElectroGalvanized Steel
 Stainless Steel 304
 Stainless Steel 316L
 Others, specify:

36. Control System

- Standard Esco Sentinel Microprocessor
 Industry Grade HMI/PLC

**37. Parameters to Monitor
(Tick All That Apply)**

- Velocity
- Pressure across filters
- Temperature
- Humidity
- Pressure in isolator
- Others, Specify:

38. Utility Requirement

- 100 VAC 50/60 Hz 1 Ph
- 115 VAC 50/60 Hz 1 Ph
- 230 VAC 50/60 Hz 1 Ph
- 380 – 400 VAC 50/60 Hz 3 Ph
- 480 VAC 60 Hz 3 Ph
- Other:

39. Optional accessories

- Electrical outlets, indicate the Type Code and Power/
Current Rating Required:
- N₂
- Drain Connection
- WFI/PW
- Compressed Air
- Exhaust Duct Connection
- Others, specify:

- Network Connections
- Adjustable Hydraulic Stand
- BioVap™ Bio-decontamination System
- CCTV Integration + Provision
- Continuous Liner System
- Glove Leak Tester
- Particle Counter (Viable/Non-viable)
- Rear View Screen Adaptation or External Monitor and
Keyboard (connected at the side with a mouse arm)
- Side Exhaust Connection
- Others, please specify:

**40. Validation
Documentation**

- FAT Protocols
- SAT Protocols
- IQ/OQ Protocol
- Biodecon Cycle Development
- Cleaning Coverage Validation
- Others, please specify:

41. Site Services

- Full Installation
- Commissioning
- Site Acceptance Test (SAT)
- User and Service Training
- Installation Qualification (IQ)
- Operational Qualification (OQ)
- Preventive Maintenance (PM)

(If required, we will provide a proposal for travel cost and daily rate)

Important: Save the completed PDF form (use menu File - Save).