

Form No.: LPF013

Version: 01

Effective Date: OCT 0 8 2020

## Sample Submission—Test Request

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Section 1: Information of Requestor for Sample Analysis (Only One Sample per Form)			
Client/Company Name:			
Client/Company Address:			
Contact Name and Designation:			
Contact E-Mail and Phone:			
Investigation/deviation Communication: ( Please provide contact name, company and email)			
Section II: Sample Information			
Sample Description:			
Manufacturer:			
Manufacturer Lot #:	Manufacturer Expiration:		
Client Lot #: Client Receiving #:			
Sample Form (check): □ Powder □ Tablet □ Capsule □ Other (list):			
Sample DEA Schedule (check): $\square$ N/A $\square$ I $\square$ III $\square$ IIIN $\square$ IIIN $\square$ IV $\square$ V			
Number of Containers: Quantity/Container:			
Sample Manufacturing Phase (check):			
□ Raw Material □ In-Process □ Finished Product □ Commercial □ Stability □ Development □ Validation □ Other (list):			
Sample Storage Condition(s):			
Section III: Tests Requested			
Test/Analysis name	Method Reference	Specification/Protocol Reference	Additional Information
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Note: For full testing, list the specification number. Fill one form for each sample. Send the original form and keep the copy for records.			
Section IV: Sample Receiving Record (For Liberty Pharma Inc. Use Only)			
Sample Received By/Date:			
Carrier Name: Airway/Shipping #:			
Additional Notes:			
Liberty Pharma Inc. Assigned AR No. (tracking purposes):			
Head-QU/Designee Approval/Date:			
Notes: Completed form filed with QU. Copy of the completed form to be filed with the completed sample records. These services are governed by Liberty Pharma Inc.'s terms and conditions unless a Master Quality/Services Agreement between the parties is in place, in which case that Master Quality/Services Agreement			
governs the services.			