This page and all fields are REQUIRED with any sample(s) submitted. Incomplete forms will result in delays. For Internal Cambrex Use Only

Cambrex
Carriorex

Send Samples with Paperwork To:

Cambrex Attn: Sample Management 3501 Tricenter Blvd, Suite C Durham, NC 27713

Print Date:

DEA Registration: RA0553051

Page ____ of _

Initials and Date:

Billing / Report Information:				4D 1	
Contract #:	Final Report Information		Phase	of Development	
Cambrex Contact:	Attn:		\Box R/D	\Box GMP	
Controlled Drug Substance (if applicable)	Company:				
Schedule 1 Schedule 2 Schedule 3	3 - 5 Address:		If GM	P, Select Phase	
DEA Number:	City/St/Zip:		□ Ph	ase 1 / Phase 2	
Turnaround Time (TAT)	Phone:		□ Ph	ase 3 Commercial	
☐ Standard ☐ Expedited	Email:				
Sample Information: An SDS is required for all materials submitted.					
Qty Amount Lot #	Sample Description	Storage Condition	Testing Information	Specification	
		Select			
Sample Disposition (if not selected, Discard all Samples will be assigned)	Return		Substances (RS)	Biopharmaceutical Samples (If applicable)	
Discard all samples (30 days after report generated)	Courier: ☐ FedEx ☐ UPS		qualified as RS	BSL 1	
□ Do Not Dispose. Store per Contract.	Acct #:		Stored/Managed	Segregated Material	
	Return all samples (30 days after report generated) Return unused portions only (30 days after report generated)	as RS		BSL 2	
Additional Comments:	☐ Return unused portions only (30 days after report generated)	□ To be	used as RS		
Client Signature and Date: Submission of this form and associated samples for testir Cambrex's general terms and conditions covering these s		incorporat	ted in the Contract	, or if none,	

Cambrex

Sample Submission Form (SSF) Reference Sheet

Billing / Report Information:

- Contract #: Write in the contract number for the testing requested.
 - o (Ex. ABCD-D23-D001)
- Cambrex Contact: The Point of Contact at Cambrex for your materials.
- Controlled Drug Substance (CDS):
 - o If a CDS, select the appropriate schedule.
 - DEA Number: Add the DEA Registration Number of the Client.
 - Please also ensure a current DEA certificate is on file with Cambrex or include with the Sample.
- **Turnaround Time (TAT)**: Indicated in the quote/contract, select the appropriate Turnaround Time on the SSF.
 - Expedited testing is subject to availability and terms of the quote or contract.

Final Report Information:

• Clearly indicate who should receive the final report and any correspondence related to this project.

Project Designation:

• Indicate if testing will be GMP or R/D. If GMP is selected, indicate the appropriate group of Phase [(Phase 1 / 2) or (Phase 3 / Commercial)].

Sample Information:

- Qty: Quantity of containers for that Sample.
- Amount: Amount and units of Sample Material shipped.
- Lot #: Lot number to be reported on the Final Report.
 - O Samples received without a lot number will be reported as N/A.
- **Sample Description**: Briefly describe each Sample as needed for clear identification.
 - Examples include Sample Strength, Stability, (Beginning, Middle, End), etc.
 - Description should match the label on the container to avoid delays.
 - Indicate if the material is considered "Bulk" Reference Substance.
- Storage Condition: The appropriate condition for storage upon receipt.
 - o If not clearly indicated, Samples will be stored per shipping conditions until clarified.

- Testing Information and Specification:
 - o GMP:
 - List the specific Client or Cambrex Test Method(s), Protocol(s) or Specification(s).
 - If Cambrex, must us the following: DUR-QCT-xxxx-TM, DUR-T-xxxx-PROT, DAV1000xxx, ect.
 - If Client, include attachment with SSF.
 - Ensure information provided is current.
 - List specific Compendial Method(s).
 - If none of the above is provided, Testing <u>WILL</u> be delayed.

o **R&D**:

- Specify Development test(s).
- List Report Results or specification.

Sample Disposition

- Select the appropriate disposition for your Sample(s).
- If return, add FEDEX or UPS account #. If not added, a pass-through cost will be applicable.

Reference Substances (RS)

- To be qualified as RS: The material being sent is for reference substance qualification testing, a COA or SOA will be generated, and may be used as a reference substance.
- **To be Stored/Managed as RS**: Projects focused on RS management, distribution, and qualification.
- To be used as RS: The material is to be used within routine and development workflows.

Biopharmaceutical Samples

- Select the appropriate Bio Safety Level (BSL). Refer to Cambrex Contact if unknown.
 - o Cambrex does not accept BSL 3 or 4.

Additional Comments:

• Use this section to document any additional information / instruction.

Separate SSF Needed for the Scenarios below (not all-inclusive):

- Separate Quote/Contract #'s.
- Multiple CDS Schedules.
- Different Turnaround Times.
- Both R/D and GMP Samples are being submitted.
- If additional testing is requested.