

SAMPLE SUBMISSION FORM

Reference SRP-13 (ROC), QA-3 (SLC)

Quote/ MSA #		PO#		[Sample Disposition	□ DO NOT OPEN □ Return / Hold fo □ Discard			
Contact:	0:			Billing Inform Company: Contact: Address: Email:	nation: - - - -				
Phone:				_ Phone:	-				
Sample Information:							_		
Test Name or Code (AAMI, ISO, US		Test Article	Name		Part #	Lot #		nples d Samples No. of Samples to be Tested	Sample Status
									☐ GLP☐ STAT☐ RES
									☐ GLP☐ STAT☐ RES
									☐ GLP☐ STAT☐ RES
									☐ GLP☐ STAT☐ RES
									☐ GLP☐ STAT☐ RES
Replacement Value:									
Comments or Special	Handling Instruct	ions:					☐ Samp	le Pooling Ins	tructions
Submitted Material Type	oe: Medica	l Device		naceutical	Other:				
Storage Conditions:	Conditions: ☐ Room temperature ☐ Refrig			erate (2-8°C)					
Applicable Departme	nts: (please mai	k all that app	ly)						
☐ Packaging: Comp	lete Section A (S	ponsor Auth.)	☐ Chemistr	y: Complete S	Section A (Sponso	or Auth)		
☐ Microbiolog y / BI (Sponsor Auth.)	/ Reusables: Co	nplete Sectio	n A		tion A (Spons	ology/Biocompatib sor Authorization) a D)			

Section A: Sponsor Authorization for testing, sign/date below:

Reports will be released	by email unless	otherwise reques	ted.		
				Internal Use On	nly
Sponsor Authorization		Date	Received By		Date
Unless other arrangement terms are net 30, prices ar for 6 months after acceptar of Be Completed if	e FOB Canyon Lance of the quote. Sample Retu	abs and are valid	Approved By Test Number(s): Test numbers veri GLP Study No(s): (if applicable)	fied by: Initial/date	Date
☐ Same as above	1				
ddress 			0		
ity			State/Province:		
ip/Post Code			Country:		
hone					
hipping Conditions	☐ Ambient/ F	toom Temp [☐ Ice Packs ☐ Dry	Ice	ontainers
hipping Information nclude account number)	☐ FedEx Acco	unt:	UPS Account:	☐ Other:	
hipping Speed	☐ Next Day	☐ Next Day - Earl	y 🗌 2 Day 📗 Gr	ound	
pecial Instructions	☐ Return Da	ta Loggers Hole	d for Pick-up 🔲 Oth	ner:	
1. Label every sample of the comprocessed until we have a comprome	for quick identife of of your Sample of your Sample of your Sample to ensure the shipped ting. If your samon the shipmen	ication. Please lak le Submission Fo d Sample Submis ire your samples § in overpacks. Ple ples are GLP, nee	rm with your shipm sion Form. get here safe. If wor ase label the overp	ent. Testing cannot king with the packa	begin nor samples ging test lab, it is we know to
5. Ship samples to the			Location:	San Diego, CA	Location:
Salt Lake City, UT Location: Attn: (Your Quote Number) Canyon Labs 16217 S Bringhurst Dr Suite 600		Attn: (Your Quote Number) Canyon Labs 7500 W Henrietta Rd Rush,		Attn: (Your Quote Number) Canyon Labs 2865 Scott St. #103 Vista, CA 92081	

NY 14543

Bluffdale, UT 84065

Addendum A

Section B: Toxicology and/or Biocompatibility Specific Test Requirements (to be completed where applicable):

Note: if testing requires a formal protocol (as requested by sponsor), the approved protocol will supersede information selected on this form.

Have samples been submitted sterile? * *Note: Samples that are sold and used sterile should be submitt testing.	ed sterile or be processed by the intended sterilization method prior to
☐ Yes, customer sterilized per following method: ☐ Ethylene Other:	Oxide
☐ No, Canyon Labs to process per following method prior to to Other:	testing: Ethylene Oxide Steam
☐ No, samples not intended to be sold or used sterile.	
☐ N/A (Testing to be performed on sample as submitted)	
Submitted Material Type:	Safety Precautions:
☐ Plastic/polymer	☐ None/unknown (standard precautions will be used)
□ Elastomer	☐ SDS enclosed
☐ Coated material, composite, laminate, metal	Flammable
☐ Pharmaceuticals	☐ Biohazard level
Other	Other
☐ Test entire sample, sample may be subdivided/cut into app	ropriate sizes for testing.
☐ Test entire sample, DO NOT sub-divide (cut) test sample for	•
☐ Do NOT test entire sample (Identify specific components of	
If Samples are submitted for testing requiring USP/ISO big	ocompatibility-based extractions, complete section B, where
applicable below.	ocompatibility-based extractions, complete section B, where
	sed testing requiring direct application, complete section C,
where applicable on Page 3.	
Canyon Labs Biocompatibility Questionnaire (for ISO 109) Test (10993-18) Questionnaire may be required for protoc	93 GLP studies) and/or Canyon Labs Chemical Characterization of preparation.
ion C: Extraction Method Selections: ☐ N/A OR ☐ as se	lected below
Extraction Methods for GPMT, Irritation Test, Systemic Injury	ection Test, Material Mediated Pyrogen, Hemolysis Tests
Extraction Condition Options:	
☐ 121°C – 1 hour ☐ 70°C – 24 hour ☐ 50°C – 72 ho	ur ☐ 37°C – 72 hour (requires justification) ☐ Other:
 Extraction Media Options (unless otherwise requested 	by Sponsor):
☐ Saline & Vegetable Oil (GPMT, Irritation, and Systemic	Toxicity tests, unless indicated in "Other" below)
☐ Saline (Material Mediated Pyrogen test)	
☐ Saline, Vegetable Oil, 1:20 Alcohol:Saline, & Polyethyle	
☐ Phosphate Buffered Saline (PBS) (ASTM Hemolysis Te	ne Glycol (USP <88> Class VI plastics testing)
☐ Other:	
	st)
Extraction Methods for Elution Cytotoxicity Test (USP <87	st)
Extraction Methods for Elution Cytotoxicity Test (USP <87 • Extraction Condition/Media Options:	st) '> or ISO 10993-5):
Extraction Methods for Elution Cytotoxicity Test (USP <87 Extraction Condition/Media Options: 37°C – 24 hour in Serum Supplemented MEM (for device)	r> or ISO 10993-5): tes with contact < 24 hours)
Extraction Methods for Elution Cytotoxicity Test (USP <87) Extraction Condition/Media Options: 37°C – 24 hour in Serum Supplemented MEM (for device of the condition) and the condition of the	st) Z> or ISO 10993-5): Des with contact < 24 hours) Des with contact ≥ 24 hours)
Extraction Methods for Elution Cytotoxicity Test (USP <87 • Extraction Condition/Media Options: □ 37°C – 24 hour in Serum Supplemented MEM (for device)	st) Z'> or ISO 10993-5): Des with contact < 24 hours) Des with contact ≥ 24 hours)

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L	Extraction Methods for BET/Rabbit Pyrogen test (Lot Release):
	Extraction Condition/Media Options
	☐ As per Test System Suitability (Method Validation): Reference Validation Study #:
	As per Canyon Labs internal work instruction, PYT-8
Tox	icology and/or Biocompatibility Specific Test Requirements (Continued from page 2; to be completed where applicable):
Sec	tion D: Direct Contact Method Selections: N/A OR as selected below
	☐ Cytotoxicity (Direct Contact Test)
	☐ Sensitization (Closed Patch Test)
Ī	☐ Skin Irritation Test
	☐ ASTM Hemolysis
	☐ Implantation Evaluation:** ☐ Histopathology (required per ISO 10993-6) ☐ Macroscopic (USP <88>)
	**Samples for histology and post mortem clinical evaluation will be submitted to an approved vendor. Samples submitted for implantation testing must be submitted sterile or be sterilized before testing (make appropriate selection above).