Libertyville - Dose Audit Test Request Form



Ship samples and completed form to: STERIS • 1880 Industrial Drive Libertyville, IL 60048 • (847) 367-5110

Company: Company name as it appears on PO or STERIS Assignment of Responsibility, this name will appear on reports			
Customer Information			
		Bill to Company:	
	•	PO / Last 4 Credit Card #:	
	Contact Person:	Quotation # (if known):	
Ι	Phone:	E-mail Report to: Provide email address to send reports to	
	Sample Description /		
on on	Part #: Information entered here should match product markings and will appear on reports		
Product Information	Unique Identifier		
Pro orn	(Lot, Run #, etc.): Information entered here should match product markings and will appear on reports as entered		
Inf	Product Registration (Medical Device, Tissue, Cosmetic, etc.): Information entered here will appear on reports and is necessary for irradiation requests		
	Samples contain hazardous material. To prevent testing delays, please attach SDS to this request. A selection must be made here: Yes No Modality: Ebeam Gamma X-ray If transferring from one radiation modality to another refer to ISO 11137 for requirements		
Processing		·	
	Verification Dose Range: kGy to kGy This is needed if sublethal irradiation is requested for Sterility samples	Shipping Carrier: This information will be used if the Lab is sending samples for irradiation as part of a transfer or routine QDA and processing is	
	This is needed if subjection in addition is requested for Sternity samples	NOT going to be completed at the RTC	
	Minimum Routine Processing Dose: kGy to kGy	Account #:	
	This is needed if high dose irradiation is requested for method validations		
P	Note: Provide additional sample to radiation facility for mini-mapping	Priority:	
Lab Transfer	Bioburden - ID up to 3 isolates (see below to opt out)	Sterility	
	Complete this box only if RE testing is requested	Complete this box only if B&F testing is requested	
	Method Validation-Inoculated (Preferred)	☐ Method Validation (1 media), 3 samples Aerobic testing	
	Method Validation-Exhaustive (Native) Bioburden Sample Quantity: 10 required selection must be made:	Method Validation (2 media), 6 samples Aerobic + Anaerobic testing Sterility Sample Quantity: □ 10 □ 100 □ other:	
Testing Information	select recovery type(s) Aerobe Fungi Anaerobe Sporeformer	Sterinty Sample Quantity. 10 100 100 other:	
	Product Description #: STERIS PD number, or Validation report number		
Te	Alert Limit:	Product Description #: STERIS PD number, or Validation report number	
I	Action Limit:		
	Check all that apply. Provide details in "Notes" below. (Additional charges may apply)		
	STAT processing requested		
ns	Refrigerate samples prior to testing		
ctio	Freeze samples prior to testing		
ıru	Do not perform Org. ID		
Inst	Pooled results		
ial	☐ Samples contain human allograft tissue ☐ Provide bacterial endotoxin testing. Quantity: if LAL is requested please indicate Turbidimetric or Chromogenic and qty		
Special Instructions		• • • • • • • • • • • • • • • • • • • •	
	☐ Do not cut or destroy samples during testing ☐ Return test samples	Ship to: Carrier:	
	Return extra samples	Account #:	
	(Provide shipping information to the right)	Priority:	
tes	Indicate any special instruction	111011031	
Notes			
Te a	By signing and dating below, I acknowledge I have read, understand and accept STERIS Terms and Conditions		
Approval	-		
	Signature (testing cannot proceed without signature and date) Date		
STERIS	Date Received:	TRFID#:	
STI O.	Received By:		

WI-01756 Attach: D Rev: 8 Eff Date: Nov 24, 2020 Status: 07e. Completed: Page 1 of 2

Multiple or Single Facility

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All microbial identification will be sourced to STERIS AST Laboratory Brooklyn Park, or other ISO 17025 accredited laboratory; however, Customer may request microbial identification samples to be sent to an alternative external laboratory service provider. Please indicate alternative site for external laboratory testing in the Notes above.

WI-01756 Attach: D Rev: 8 Eff Date: Nov 24, 2020 Status: 07e. Completed: Page 2 of 2

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