

	Report #:
	COA261129-013
	Page 1 of 1
Title: Certificate of Analysis (CoA)	
Bacterial Endotoxins Test (ABI- TEST 0012)	

Date:
Date Tested:
Customer:
Testing material:
Lot Number:
Sample ID:
Content/Potency:
Storage:
Packing material:
Labeled as:
Manufacturer:
Testing Purpose:

8/20/2026
8/15/2026
Finnrick
Retatrutide
Y6ST6UG
BTL261129-013
10 mg
R.T.
Small clear vial: white sample, white label, silver crimp, white plastic cap.
Retatrutide
N/A
(ABI-TEST 0012) test quantifies bacterial endotoxin levels to confirm that the product does not elicit pyrogenic reactions, maintaining patient safety and product integrity.



Test	Method	Specification	Result
Bacterial Endotoxins Test	USP <85>	350 EU/vial reconstituted	PASS

The results of the CoA relate only to the item(s) tested and applied to the sample as received.

Bacterial Endotoxins Test Retatrutide Lot Y6ST6UG

Sample ID	Description	Volume (mL)	Endotoxin Limit (EU/vial)	Lysate Sensitivity (λ, EU/mL)	Result
Y6ST6UG	Retatrutide	2	350 EU/vial	0.05	4.5 EU/vial

The sample passed a valid kinetic turbidity endotoxin test based on the type, mode of administration and dosage.
Endotoxin Limits is established per USP <85> guidelines for injectables

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