BTLabs

Report #: IF-824-QSA-TR008-2026-094-03

Page 1 of 1

Title:

Certificate of Analysis (CoA)

Bacterial Endotoxins Test (ABI- TEST 0012)

Date: 2/21/2026
Date Tested: 2/19/2026
Customer: Finnrick
Testing material: Tesamorelin
Lot Number: YV5MGBT
BT Sample ID: 005000038919434
Labeled Peptide Content/Potency: 20 mg
Storage: R.T.
Packing material: small clear vial: white sample, white label, bright red crimp, red plastic cap.
Labeled as: Tesamorelin
Manufacturer: N/A
Testing Purpose: (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>	40 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 Tesamorelin Lot YV5MGBT

Sample ID	Description	Volume (mL)	Endotoxin Limit (EU/vial)	Lysate Sensitivity (λ, EU/mL)	Result
005000038919434	Tesamorelin	3	40 EU/vial	0.25	15.8 EU/vial

The sample passed a valid gel clot endotoxin test based on the type, mode of administration and dosage.

Andrea Castro, AS

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Report #: IF-824-QSA-TR006-2026-94-80

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Title:

Certificate of Analysis (CoA)

Date: 3/02/2026
Date Tested: 3/02/2026
Customer: Finnrick
Testing material: Tesamorelin
Lot Number: YV5MGBT
BT Sample ID: 005000038919434
Labeled Peptide Content/Potency: 20 mg
Storage: R.T.
Packing material: small clear vial: white sample, white label, bright red crimp, red plastic cap.
Labeled as: Tesamorelin
Manufacturer: N/A
Testing Purpose: FTIR and HPLC analysis for the identification, purity, potency and composition of a peptide product. It does not provide information on particulate matter, microbial contamination or presence of endotoxins.

Test	Method	Specification	Result
General Appearance	USP <630>	white powder	white powder
Mass	USP <41>	As recorded	65.6 mg
FTIR Identification and Composition Analysis	USP <197A>	Sample spectrum should confirm the content of peptide via characteristic bands	FTIR sample spectrum confirms the presence of Tesamorelin with addition of excipient(s)/fillers.
HPLC Purity of Peptide Assay	USP <621>	Specifications: ≥ 98 %	99.8 %
HPLC Potency Assay	USP <621>	Specifications: 95 – 105% of 20 mg	19.4 mg (97.3 %)
Peptide-to-Excipients Ratio	USP <1151>	Specifications: Ratios of 1:2 to 1:10 peptide: excipients	19.4 : 46.2 mg (1:2.4)

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

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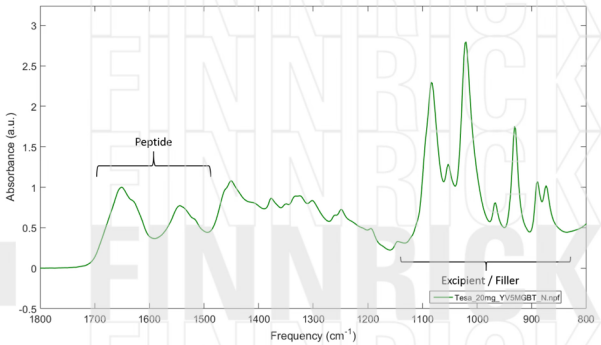
Report #:
IF-824-QSA-TR006-2026-94-80
Page 2 of 2

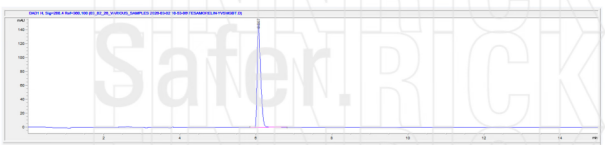
Title:

Certificate of Analysis (CoA)

FTIR ID and Composition Analysis: Tesamorelin Lot YV5MGBT



HPLC Purity and Potency Assay @ 280 nm: Tesamorelin Lot YV5MGBT



Tesamorelin Lot YV5MGBT @ 280 nm		
Peak #:	Retention Time (min)	Area (mAU*s)
1	6.067	532.1

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