

Customer:
Research Labs

EA Sample ID: 26EA0814-026
Sample Name: Tesamorelin
Sample Type: Powder
Batch/Lot: TSA10-0726-6

Date Received:
08/14/2026
Date Completed:
08/17/2026



ETHOS
ANALYTICS

CERTIFICATE OF ANALYSIS

Summary of Results

<u>Analysis Type</u>	<u>Method</u>	<u>Date Tested</u>	<u>Status</u>
Tesamorelin	HPLC	08/17/2026	Complete
Bacterial Endotoxins	USP <85>	08/17/2026	Complete
Heavy Metals	ICP-MS	08/16/2026	Pass



Serving Size: N/A

Label Claim Profile

<u>Analyte</u>	<u>Result</u>	<u>Specification</u>	<u>LOQ (%)</u>
Tesamorelin	10.26mg/vial	10mg/vial	0.1
Identity of the analyte was confirmed by retention time concordance with a certified reference standard, per USP <621>.			
Chromatographic Purity	>99%	>99%	0.1
Bacterial Endotoxins (Gel-Clot: 1:10 Dilution)	<1.25 EU/ml	<5 EU/ml	$\lambda =$ 0.125 EU/mL

NOTES: Injectable peptides may include acceptable salts, sugars, or buffering agents added to improve solubility and stability. These excipients are non-chromophoric, typically not detected by HPLC, and are not considered impurities.

LOD = LIMIT OF DETECTION; LOQ = LIMIT OF QUANTIFICATION



Ethos Analytics Laboratory
3020 E Camelback Rd STE 397
Phoenix, AZ 85016
Info@Ethosanalytics.io
www.Ethosanalytics.io
Lic #: 000026LRCND60176649
ISO/IEC 17025 Acc #: 117798

Noel Samsum
Laboratory Director
17-Aug-2026

The sample analyzed was inspected and is free from visual mold, mildew, and foreign matter. The testing procedures, equipment calibration, and maintenance are all in accordance with ISO/IEC 17025:2017 standards. The presented report is only applicable to the sample specified above and may not be applied to any similar or identical products. Reports are prohibited from being reproduced with alterations of any kind.

Customer:
Research Labs

EA Sample ID: 26EA0814-026
Sample Name: Tesamorelin
Sample Type: Powder
Batch/Lot: TSA10-0726-6

Date Received:
08/14/2026
Date Completed:
08/17/2026



ETHOS
ANALYTICS

CERTIFICATE OF ANALYSIS

Heavy Metal Analysis (USP <233>)

<u>Analyte</u>	<u>Result</u> (mcg/vial)	<u>LOQ (ppm)</u>	<u>LOD (ppm)</u>	<u>Limit</u> (mcg/day)	<u>Pass/Fail</u>
Arsenic	0.002	0.010	0.005	0.5	Pass
Cadmium	<LOQ	0.010	0.005	0.1	Pass
Lead	0.003	0.010	0.005	0.5	Pass
Mercury	<LOQ	0.010	0.005	0.1	Pass

Microbiological Analysis (USP <61>, USP <62>)

<u>Microbe</u>	<u>Result</u>	<u>Limit</u>	<u>Pass/Fail</u>
Total Aerobic Plate	Not Tested	<100 CFU/g	-
Coliform	Not Tested	<10 CFU/g	-
Bile-Tolerant Gram-Negative Bacteria	Not Tested	<10 CFU/g	-
Yeast and Mold	Not Tested	<10 CFU/g	-
Escherichia Coli	Not Tested	Negative/1g	-
Salmonella	Not Tested	Negative/1g	-
Staphylococcus Aureus	Not Tested	Negative/1g	-

NOTES:

CFU = Colony Forming Unit
NS = Not Specified
NT = Not Tested

LOQ = Limit of Quantification
LOD = Limit of Detection



Ethos Analytics Laboratory
3020 E Camelback Rd STE 397
Phoenix, AZ 85016
Info@Ethosanalytics.io
www.Ethosanalytics.io
Lic #: 000026LRCND60176649
ISO/IEC 17025 Acc #: 117798

Noel Samsum
Laboratory Director
17-Aug-2026

The sample analyzed was inspected and is free from visual mold, mildew, and foreign matter. The testing procedures, equipment calibration, and maintenance are all in accordance with ISO/IEC 17025:2017 standards. The presented report is only applicable to the sample specified above and may not be applied to any similar or identical products. Reports are prohibited from being reproduced with alterations of any kind.