

## Analysis Report

Official laboratory analysis summary for the submitted sample and associated quality-control review.

SAMPLE <b>Pinealon</b>	RECEIVED DATE <b>Mar 25, 2026</b>	ANALYSIS DATE <b>Mar 27, 2026</b>	REPORT GENERATED <b>Mar 30, 2026</b>
STRENGTH	<b>20mg</b>	MANUFACTURER	<b>PepticoresAminos</b>
BATCH NUMBER	<b>PC-PIN20-0130U</b>	LAB CODE	<b>989-1</b>
CLIENT	<b>www.pepticoresaminos.net</b>		

## SAMPLE INFORMATION

## Pinealon

**20MG**FORM **White powder in a glass vial**SAMPLE SUBMISSION **Sample provided by customer**BATCH **PC-PIN20-0130U**CAP / CRIMP COLOR **green/black**RECEIVED DATE **Mar 25, 2026**

SAMPLE IMAGE

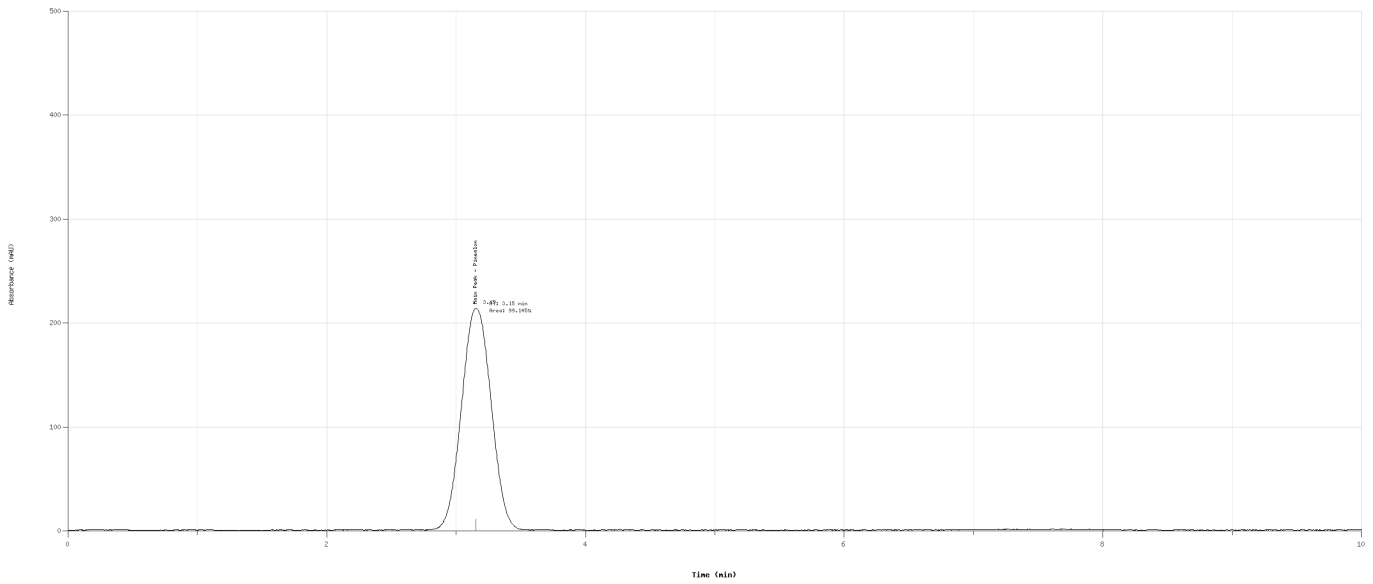
## ANALYTICAL SUMMARY

IDENTITY	<b>Pinealon</b>
PURITY	<b>99.145%</b>
QUANTITY	<b>19.9mg</b>
BATCH	<b>PC-PIN20-0130U</b>
MANUFACTURER	<b>PepticoresAminos</b>

# RP-HPLC-UV CHROMATOGRAM (220 NM)

Detection: UV 220 nm

Sample ID: #lineation  
Report ID: 578-2020-094005  
Method: RP-HPLC-UV Short Peptide  
Detector: UV 220 nm | Runtime: 12.0 min



## METHOD

TIME	H2O + 0.1% TFA	ACN + 0.1% TFA
0min	96%	4%
2min	96%	4%
10min	12%	88%
12min	12%	88%

## TECHNICAL NOTE

Gradient RP-HPLC-UV screening method with UV detection at 220 nm. Runtime 12.0 min.

## COMMENTS

Results indicate that the sample conforms to the expected specifications with no unusual findings.

## ANALYSIS & METHODOLOGY

### STANDARD PEPTIDE PURITY, MASS & ADDITIONAL TESTING

HPLC-UV: High-Performance Liquid Chromatography with Ultraviolet detection. Additional testing includes bioburden screening, endotoxin analysis and heavy metals assessment.

## BIOBURDEN

TEST	RESULT	UNIT	REPORTING LIMIT
<b>Total Aerobic Microbial Count</b> USP <61>/Eur. Ph. 2.6.12. Plate Count Method	Not detected	CFU/g	>= 1000
<b>Total Yeast and Mold Count</b> USP <61>/Eur. Ph. 2.6.12. Plate Count Method	Not detected	CFU/g	>= 100

## ENDOTOXIN ANALYSIS

TEST	RESULT	UNIT	REPORTING LIMIT
<b>Bacterial Endotoxin</b> USP<85>/ Eur. Ph. 2.6.14. Bacterial Endotoxin Chromogenic Test	< 0.001	EU/mg	> 0.5

## HEAVY METALS

TEST	RESULT	UNIT	REPORTING LIMIT
<b>Arsenic</b> Elemental Impurities Screening	Not detected	ppm	>= 1.5
<b>Cadmium</b> Elemental Impurities Screening	Not detected	ppm	>= 0.5
<b>Cobalt</b> Elemental Impurities Screening	Not detected	ppm	>= 25
<b>Lead</b> Elemental Impurities Screening	Not detected	ppm	>= 1.5
<b>Nickel</b> Elemental Impurities Screening	Not detected	ppm	>= 25
<b>Quicksilver</b> Elemental Impurities Screening	Not detected	ppm	>= 1.5
<b>Vanadium</b> Elemental Impurities Screening	Not detected	ppm	>= 25

## TECHNICAL APPENDIX

This appendix documents the analytical methodology, instrumentation and acceptance criteria applied for the evaluation of the sample.

## METHOD SPECIFICATION

PARAMETER	EARLY-ELUTING SHORT PEPTIDE RP-HPLC
<b>ANALYTICAL MODE</b>	Short peptide RP-HPLC-UV screening
<b>COLUMN</b>	C18 peptide column, short-gradient configuration
<b>MOBILE PHASE A</b>	Water + 0.1% TFA
<b>MOBILE PHASE B</b>	Acetonitrile + 0.1% TFA
<b>FLOW RATE</b>	1.0 mL/min
<b>DETECTION</b>	UV 220 nm
<b>INJECTION VOLUME</b>	10 uL
<b>RUNTIME</b>	10.0 min
<b>SAMPLE DILUENT</b>	Aqueous organic diluent compatible with RP-HPLC
<b>SAMPLE PREPARATION</b>	Direct dilution with clarification before analysis

## INSTRUMENT PLATFORM

PARAMETER	STANDARD HPLC-UV PLATFORM
<b>SYSTEM TYPE</b>	Analytical HPLC system

PARAMETER	STANDARD HPLC-UV PLATFORM
DETECTOR	UV/VIS detector
ACQUISITION	Chromatographic acquisition and integration software
REVIEW MODE	Retention-time and response-profile review
WORKFLOW NOTE	Used for routine peptide purity and identity screening

## ANALYTICAL CRITERIA

PARAMETER	ACCEPTANCE FRAMEWORK	BASIS
IDENTITY	Retention-time and profile agreement with reference expectations	Chromatographic identity review
PURITY	NLT 98.0% unless a stricter report-specific specification is declared	Integrated RP-HPLC-UV purity profile
QUANTITY	Measured content reviewed against the declared sample strength	Report-level analytical summary
BIOBURDEN	Not detected or within stated reporting limits	Microbial screening table
ENDOTOXIN	Below stated reporting limit / internal screening threshold	Endotoxin analysis table
HEAVY METALS	Below individual reporting limits where screened	Elemental impurities screening table

This appendix documents the analytical methodology, instrumentation and acceptance criteria applied for the evaluation of the sample.

## VERIFICATION

Verify this report through the official Synaptica Analytics verification page using the details below.

**Verification URL** [synaptica-labs.com/verify-report/](https://synaptica-labs.com/verify-report/)

**Report ID** SYN-2026-004885

**Verification Key** VK-MP4T-XLXA



## DIGITAL SIGNATURE

**DIGITALLY SIGNED BY:**

**Martin Saar**

Date: 2026.03.30

12:56:24 +02'00'

Director

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Analysis date: Mar 27, 2026

Report generated: Mar 30, 2026

Analytical testing performed by Synaptica Analytics -

Analytical Services Division

Synaptica Analytics  
**SYN-2026-004885**

Laboratory Analysis Report  
**VK-MP4T-XLXA**

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