Synaptica Analytics

Independent Laboratory Analysis Report

OFFICIAL ANALYTICAL DOCUMENT

REPORT ID
SYN-2026-005017

Analysis Report

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

SAMPLE	RECEIVED DATE	ANALYSIS DATE	REPORT GENERATED
Tirzepatide	Jul 30, 2026	Aug 10, 2026	Aug 12, 2026

STRENGTH	60mg	MANUFACTURER	PepticoreAminos
BATCH NUMBER	PC-TZ60-1007A	LAB CODE	989-1
CLIENT	www.pepticoreaminos.net		

SAMPLE INFORMATION

Tirzepatide

60MG

FORM	Clear reconstituted solution in a pre-filled pen
SAMPLE SUBMISSION	Sample provided by customer
BATCH	PC-TZ60-1007A
RECEIVED DATE	Jul 30, 2026



SYNAPTICA ANALYTICS

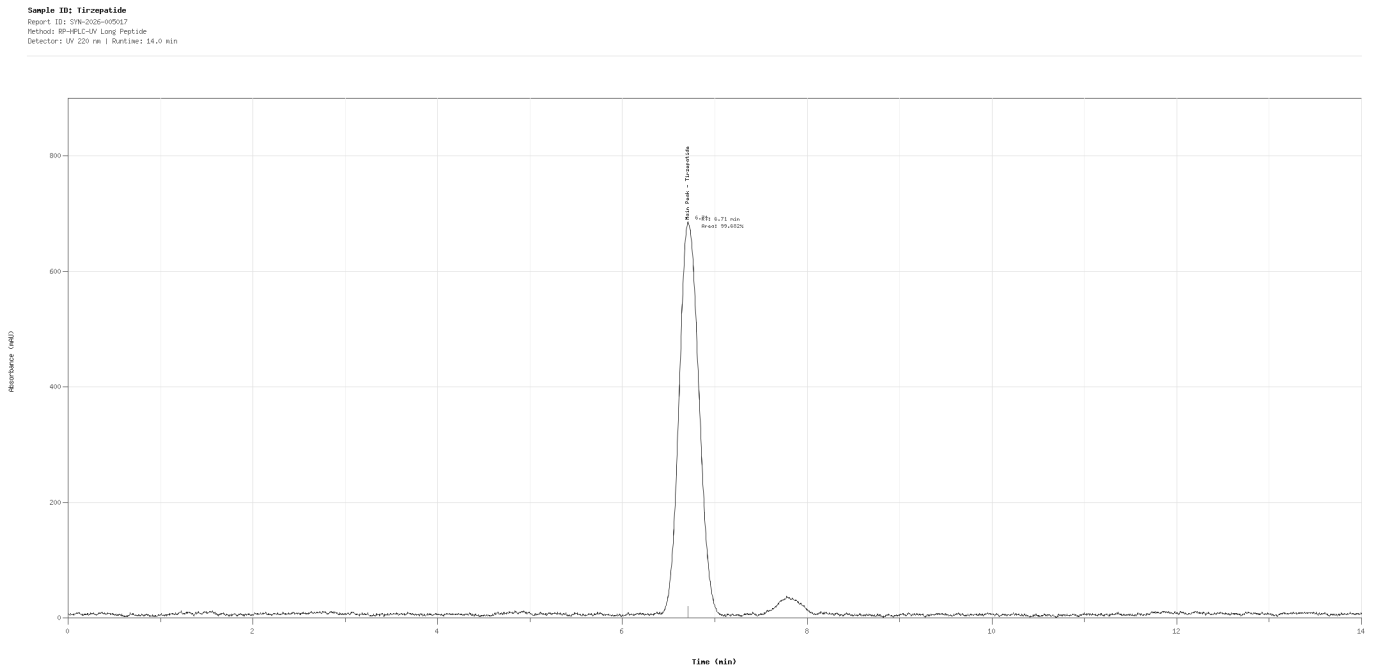
SAMPLE IMAGE

ANALYTICAL SUMMARY

IDENTITY	Tirzepatide
PURITY	99.682%
QUANTITY	60.12mg
BATCH	PC-TZ60-1007A
MANUFACTURER	PepticoreAminos

RP-HPLC-UV CHROMATOGRAM (220 NM)

Detection: UV 220 nm | Runtime: 14.0 min



METHOD

TIME	H2O + 0.1% TFA	ACN + 0.1% TFA
0min	90%	10%
2min	90%	10%
10min	8%	92%
12min	8%	92%
14min	8%	92%

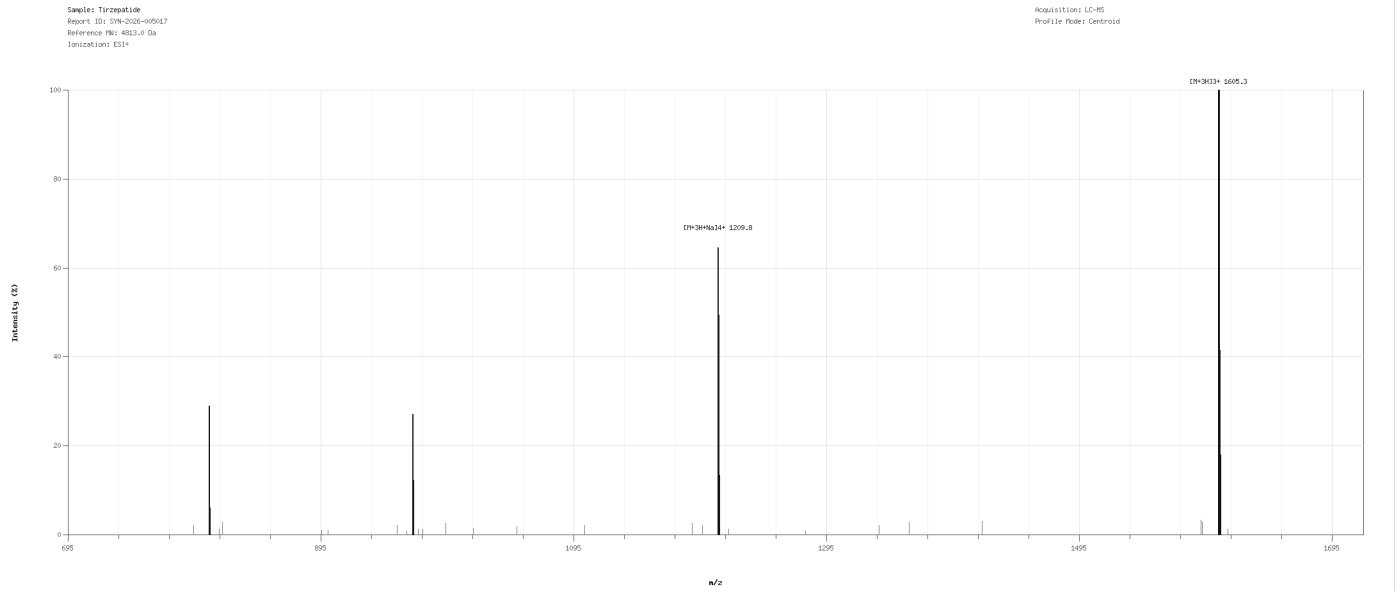
TECHNICAL NOTE

Analysis performed by RP-HPLC-UV at 220 nm using a gradient elution program. Total runtime: 14.0 minutes.

COMMENTS

The reported data support compliance of the sample with the defined analytical specifications for identity, purity and impurity profile based on the applied RP-HPLC-UV method.

LC-MS MASS SPECTRUM



Charge-state distribution and isotope clustering are consistent with the expected analytical mass response.

ANALYSIS & METHODOLOGY

STANDARD PEPTIDE PURITY, MASS & ADDITIONAL TESTING

Analysis performed using RP-HPLC-UV under validated laboratory conditions. Peak profile, retention behavior and purity response were evaluated against internal reference standards.

FINISHED PRODUCT STERILITY

PARAMETER	RESULT
Test Article	**Finished pre-filled pen**
Testing Stage	**After filling and final assembly**
Method	**USP <71> / Ph. Eur. 2.6.1 Sterility Test**
Test Type	**Direct inoculation**
Sample Size	**n = 1 finished pen**
Incubation Period	**Not less than 14 days**
Culture Media	**FTM / SCDM**
Incubation Conditions	**20-25°C and 30-35°C**
Observation	**No visible microbial growth observed**
Acceptance Criteria	**No evidence of microbial growth**
Result	**No growth observed**
Conclusion	**Passed**

ENDOTOXIN ANALYSIS

TEST	RESULT	UNIT	REPORTING LIMIT
Bacterial Endotoxin USP<85>/ Eur. Ph. 2.6.14. Bacterial Endotoxin Chromogenic Test	**< 0.005**	**EU/mL**	**NMT 0.5**

HEAVY METALS

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

COMPOUND REFERENCE

PARAMETER	TIRZEPATIDE
PUBCHEM CID	166567236
CAS	2023788-19-2
MOLECULAR FORMULA	C225H348N48O68
MOLECULAR WEIGHT	4813.0 g/mol

METHOD SPECIFICATION

PARAMETER	LONG PEPTIDE EXTENDED RP-HPLC-UV METHOD
ANALYTICAL MODE	Extended gradient RP-HPLC-UV peptide purity screen
COLUMN	C18 peptide column, extended-gradient configuration
MOBILE PHASE A	Water + 0.1% TFA
MOBILE PHASE B	Acetonitrile + 0.1% TFA
FLOW RATE	0.7 mL/min
DETECTION	UV 220 nm

PARAMETER	LONG PEPTIDE EXTENDED RP-HPLC-UV METHOD
INJECTION VOLUME	10 uL
RUNTIME	14.0 min
SAMPLE DILUENT	Water/acetonitrile compatible diluent
SAMPLE PREPARATION	Diluted to method range and clarified before injection

INSTRUMENT PLATFORM

PARAMETER	HIGH-RESOLUTION UPLC/HPLC-UV PLATFORM
SYSTEM TYPE	High-resolution LC platform
DETECTOR	UV/VIS detector
ACQUISITION	Chromatographic acquisition and integration software
REVIEW MODE	Retention-time, UV response and mass-spectrum review
WORKFLOW NOTE	Used for long peptide analytical profile review
MS DETECTOR	Mass spectrometer, ESI+ acquisition
MS ACQUISITION	LC-MS spectral acquisition and processing

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
FINISHED PRODUCT STERILITY	No evidence of microbial growth after incubation	USP <71> / Ph. Eur. 2.6.1 sterility test
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
Report ID SYN-2026-005017
Verification Key VK-PGZD-BV82

SCAN TO VERIFY

DIGITAL SIGNATURE



DIGITALLY SIGNED BY:
Martin Saar
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Analysis date: Aug 10, 2026
Report generated: Aug 12, 2026
Analytical testing performed by Synaptica Analytics - Analytical Services Division

Synaptica Analytics
SYN-2026-005017
Laboratory Analysis Report
VK-PGZD-BV82

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