

Analysis Report

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

SAMPLE GHK-Cu	RECEIVED DATE Apr 24, 2026	ANALYSIS DATE Apr 29, 2026	REPORT GENERATED Apr 30, 2026
STRENGTH	100mg	MANUFACTURER	PeptiCoreAminos
BATCH NUMBER	PC-GK100-403F	LAB CODE	987-1
CLIENT	www.peptiCoreAminos.net		

SAMPLE INFORMATION

GHK-Cu

100MG

FORM	Blue powder in a glass vial
SAMPLE SUBMISSION	Sample provided by customer
BATCH	PC-GK100-403F
CAP / CRIMP COLOR	Blue/Silver
RECEIVED DATE	Apr 24, 2026



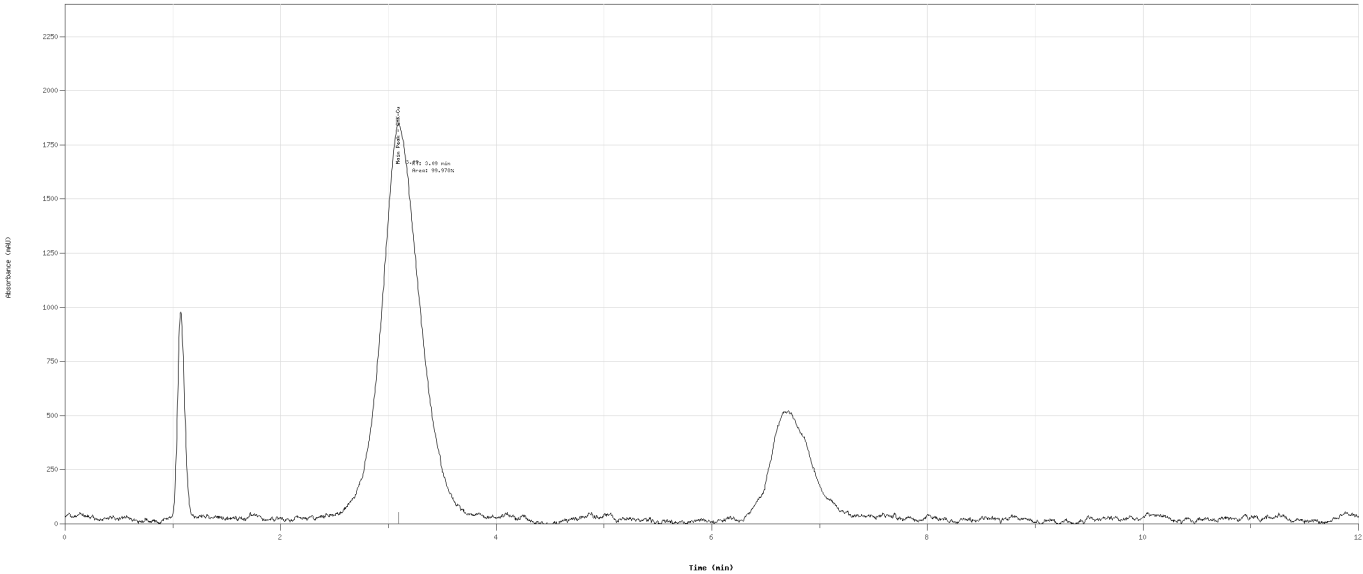
SAMPLE IMAGE

ANALYTICAL SUMMARY

IDENTITY	GHK-Cu
PURITY	99.978%
QUANTITY	102.91mg
BATCH	PC-GK100-403F
MANUFACTURER	PeptiCoreAminos

RP-HPLC-UV CHROMATOGRAM (220 NM)
Detection: UV 220 nm | Runtime: 12.0 min

Sample ID: QWK-Cu
Report ID: 519-2020-004999
Method: RP-HPLC-UV Complex Profile
Detector: UV 220 nm | Runtime: 12.0 min



METHOD

TIME	H2O + 0.1% TFA	ACN + 0.1% TFA
0min	94%	6%
2min	94%	6%
10min	18%	82%
12min	18%	82%

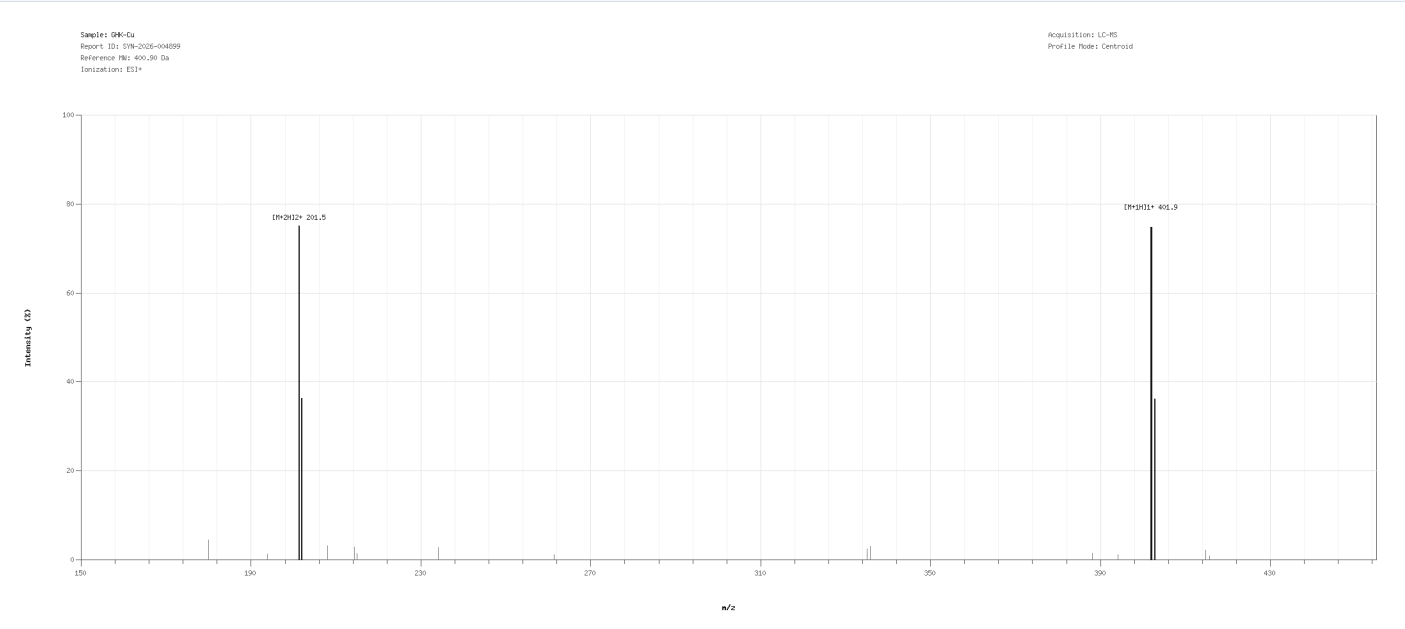
TECHNICAL NOTE

Analytical conclusions are limited to the tested sample and the conditions described in this report. Total runtime: 12.0 minutes.

COMMENTS

The sample meets the defined analytical specifications for identity, purity and impurity profile according to the applied RP-HPLC-UV analytical review.

LC-MS MASS SPECTRUM



The recorded mass spectrum shows signals compatible with the expected molecular profile of the sample.

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.

BIOBURDEN

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

ENDOTOXIN ANALYSIS

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

TEST	RESULT	UNIT	REPORTING LIMIT
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	GHK-CU
PUBCHEM CID	139035031
CAS	49557-75-7
MOLECULAR FORMULA	C14H21CuN6O4-
MOLECULAR WEIGHT	400.9 g/mol

METHOD SPECIFICATION

PARAMETER	METAL COMPLEX RP-HPLC-UV SCREENING
ANALYTICAL MODE	RP-HPLC-UV screen for metal-complex peptide profile
COLUMN	C18 compatible analytical column
MOBILE PHASE A	Water + 0.1% TFA
MOBILE PHASE B	Acetonitrile + 0.1% TFA
FLOW RATE	0.8 mL/min
DETECTION	UV 220 nm
INJECTION VOLUME	10 uL
RUNTIME	12.0 min
SAMPLE DILUENT	Aqueous organic diluent compatible with complexed peptides
SAMPLE PREPARATION	Diluted and mixed using low-metal-contact handling

INSTRUMENT PLATFORM

PARAMETER	COMPLEX-SCREENING HPLC-UV PLATFORM
SYSTEM TYPE	Analytical HPLC system
DETECTOR	UV/VIS detector
ACQUISITION	Chromatographic acquisition and integration software
REVIEW MODE	Primary peak, related profile and late-event review

PARAMETER	COMPLEX-SCREENING HPLC-UV PLATFORM
WORKFLOW NOTE	Used for specialty peptide and complex-associated profiles

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

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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-2026-004899
Verification Key VK-HQ63-MTF4

SCAN TO VERIFY



DIGITAL SIGNATURE

DIGITALLY SIGNED BY:
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Date: 2026.04.30
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Director