

Certificate of Analysis

Lab ID: **RET-426**

REPORTED September 3, 2026

CLIENT

VERIFY peptidetestcanada.ca/verify

EMAIL info@peptidetestcanada.ca

ORDER # PTC-753742

SCAN TO VERIFY



PTC-WHTQTAGU

END CUSTOMER

NAME Peptix Canada , website <https://peptixcanada.ca/>

§ 01 Sample Information

FOR RESEARCH PURPOSES ONLY

Sample ID:	RET-426	Received:	08/26/2026
Compound:	Retatrutide	Analyzed:	09/03/2026
Report No.:	COA-2026-SC-01527	CAS:	2381089-83-2
Lot Number:	PTX-027	Formula:	C₂₂₁H₃₄₂N₄₆O₆₈
Total Mass:	67.6 mg	Mol Weight:	4731.3 g/mol
Appearance:	Lyophilized powder		

SAMPLE PHOTOGRAPH



§ 02 Analytical Results

Qualitative and Quantitative chemical analysis by Liquid Chromatography Tandem Mass Spectrometry (HPLC-MS/MS)

	SPECIFICATION	RESULT	STATUS
Compound Test:	Retatrutide	Retatrutide	
Net Peptide Content:	10.00 mg	12.13 mg	
Peptide Fill Accuracy:	≥ 90 %	121.3 %	PASS
Chromatographic Purity:	≥ 98 %	99.7 %	

Peptide fill accuracy is the measured net peptide content expressed as a percentage of the declared label claim (measured ÷ label claim × 100).

Chromatographic purity is the target peptide's concentration as a percentage of the total peptide concentration measured — the target peptide against the other trace peptides detected in the same sample. It reflects peptide-related impurities only; it does not measure salts, excipients, or water, and is separate from net peptide content (the actual mg of peptide). — means not determined.

§ 03 HPLC-MS/MS Detail

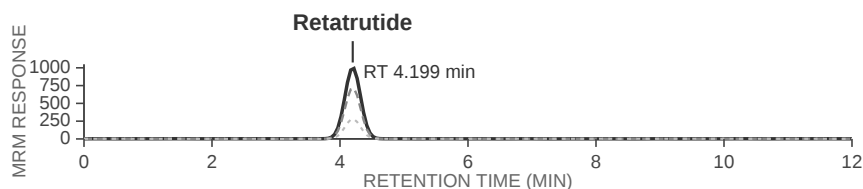
METHOD B · AGILENT 1290 INFINITY II + 6495 QQQ · ESI+

PRECURSOR [M+5H]⁵⁺

947.90 Da

Quantifier transition ● MRM

947.90 > 136.10 CE 30 V



ROLE	TRANSITION	CE
● Quantifier	947.9 > 136.1	30 V
■ Qualifier	947.9 > 110.1	35 V

Agilent Poroshell 120 EC-C18 · 2.1×100 mm · 2.7 μm · Solvent A: 0.1 % formic acid in H₂O · Solvent B: 0.1 % formic acid in Acetonitrile

DISCLAIMER · RESEARCH USE ONLY

Results relate **only to the sample as received** by Side-Chain Analytics Inc. and the tests performed, as and when received — not to any other unit, vial, lot, or batch, and not to any other product bearing the same name, label, or description. This report is not a verification of any lot or batch. **For research use only:** this report is not a certificate of quality, safety, efficacy, or regulatory compliance; the material tested is not authorized by Health Canada for human or veterinary use; and this report is not for human or veterinary administration, clinical or diagnostic use, or label claims. Side-Chain Analytics Inc. does not manufacture, source, or sell the material tested and makes **no warranty, express or implied**; its liability shall not exceed the testing fee. This report may not be reproduced except in full, and may not be reproduced, displayed, quoted, or referenced — in whole or in part, including by any badge, seal, or link — in connection with the advertising, offer for sale, sale, or distribution of any product. This certificate is valid only as currently shown at sidechainanalytics.com/verify and may be withdrawn.

Certificate of Analysis

Lab ID: **ENDO-RET-426**

REPORTED September 17, 2026
CLIENT
VERIFY peptidetestcanada.ca/verify
EMAIL info@peptidetestcanada.ca
ORDER # PTC-753742



END CUSTOMER
NAME Peptix Canada , website <https://peptixcanada.ca/>

§ 01 Sample Information

FOR RESEARCH PURPOSES ONLY

Sample ID:	ENDO-RET-426	Received:	08/26/2026
Compound:	Retatrutide	Analyzed:	09/03/2026
Report No.:	COA-2026-SC-01527-EN	CAS:	2381089-83-2
Lot Number:	PTX-027	Formula:	C₂₂₁H₃₄₂N₄₆O₆₈
Total Mass:	67.6 mg	Mol Weight:	4731.3 g/mol
Aliquot mass:	11.0 mg		
Appearance:	Lyophilized powder		

SAMPLE PHOTOGRAPH



§ 02 Endotoxin Analysis

KINETIC CHROMOGENIC LAL/TAL ENDOTOXIN TEST (USP <85>)

BACTERIAL ENDOTOXIN · KINETIC CHROMOGENIC (QUANTITATIVE)



0.19 EU/mL

Measured endotoxin 0.19 EU/mL is **within** the 1 EU/mL limit.

Expressed at 1 mg/mL reconstitution (0.19 EU/mg).

Test method	Kinetic Chromogenic LAL/TAL Endotoxin Test (USP <85>)
Measured endotoxin	0.19 EU/mL · 0.19 EU/mg at 1 mg/mL reconstitution
Positive product control	65% recovery · acceptance 50 – 200% (USP <85>)
Endotoxin limit	1 EU/mL

The endotoxin limit is established by the endotoxin test kit manufacturer. The measured value is the endotoxin burden per milligram of peptide; EU/mL and EU/mg are numerically equal at 1 mg/mL reconstitution.

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REVISION 1 Supersedes COA-2026-SC-01527-EN issued September 10, 2026. The measured value is unchanged. The previous certificate reported it as "below 0.19 EU/mL", which presented a quantified measurement as a non-detect; positive product control recovery has been added.