Peptide Test Canada
peptidetestcanada.ca
Canada's Most Trusted Independent Peptide Testing Lab

Certificate of Analysis

SCAN TO VERIFY
PTC-K8UGGDKS

Lab ID: **RET-427**

REPORTED September 3, 2026

CLIENT

VERIFY peptidetestcanada.ca/verify
EMAIL info@peptidetestcanada.ca
ORDER # PTC-753742

END CUSTOMER

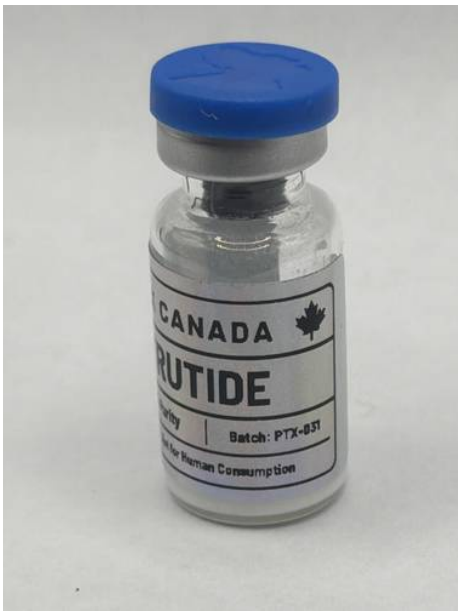
NAME Peptix Canada , website https://peptixcanada.ca/

§ 01 Sample Information

FOR RESEARCH PURPOSES ONLY

Sample ID:	**RET-427**	Received:	**08/26/2026**
Compound:	**Retatrutide**	Analyzed:	**09/03/2026**
Report No.:	**COA-2026-SC-01525**	CAS:	**2381089-83-2**
Lot Number:	**PTX-031**	Formula:	**$C_{221}H_{342}N_{46}O_{68}$**
Total Mass:	**79.0 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:	20.00 mg	**22.76 mg**	
Peptide Fill Accuracy:	≥ 90 %	**113.8 %**	**PASS**
Chromatographic Purity:	≥ 98 %	**99.2 %**	

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

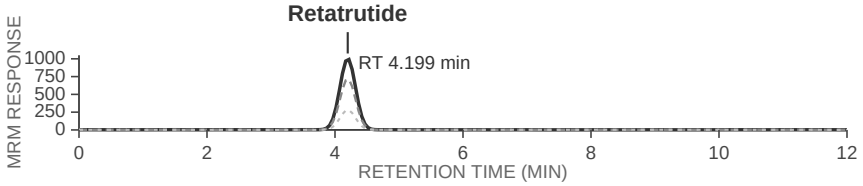
METHOD B · AGILENT 1290 INFINITY II + 6495 QQQ · ESI+

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 H_2O · Solvent B: 0.1 % formic acid in Acetonitrile

Issued September 3, 2026

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.

Issued September 3, 2026

Peptide Test Canada
peptidetestcanada.ca
Canada's Most Trusted Independent Peptide Testing Lab

Certificate of Analysis

Lab ID: **ENDO-RET-427**

REPORTED September 17, 2026

CLIENT

VERIFY peptidetestcanada.ca/verify

EMAIL info@peptidetestcanada.ca

ORDER # PTC-753742

SCAN TO VERIFY

PTC-4VBF9XBG

END CUSTOMER

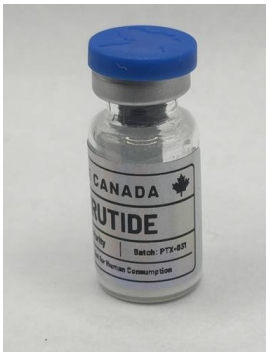
NAME Peptix Canada , website https://peptixcanada.ca/

§ 01 Sample Information

FOR RESEARCH PURPOSES ONLY

Sample ID:	**ENDO-RET-427**	Received:	**08/26/2026**
Compound:	**Retatrutide**	Analyzed:	**09/03/2026**
Report No.:	**COA-2026-SC-01525-EN**	CAS:	**2381089-83-2**
Lot Number:	**PTX-031**	Formula:	**$C_{221}H_{342}N_{46}O_{68}$**
Total Mass:	**79.0 mg**	Mol Weight:	**4731.3 g/mol**
Aliquot mass:	**10.2 mg**		
Appearance:	**Lyophilized powder**		

SAMPLE PHOTOGRAPH



§ 02 Endotoxin Analysis

KINETIC CHROMOGENIC LAL/TAL ENDOTOXIN TEST (USP ‹85›)

BACTERIAL ENDOTOXIN · KINETIC CHROMOGENIC (QUANTITATIVE)

✓

0.35 EU/mL

Measured endotoxin 0.35 EU/mL is **within** the 1 EU/mL limit.

Expressed at 1 mg/mL reconstitution (0.35 EU/mg).

Test method	Kinetic Chromogenic LAL/TAL Endotoxin Test (USP ‹85›)
Measured endotoxin	0.35 EU/mL · 0.35 EU/mg at 1 mg/mL reconstitution
Positive product control	51% 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.

REVISION 1 Supersedes COA-2026-SC-01525-EN issued September 10, 2026. The measured value is unchanged. The previous certificate reported it as "below 0.35 EU/mL", which presented a quantified measurement as a non-detect; positive product control recovery has been added.