

CJC-1295 +
Ipamorelin Blend

QUALITY CONTROL CERTIFICATE

CJC-1295 + Ipamorelin acetate blend



Certificate No.
CP-QCC-2026-055952

Lot Number
CJCIPM-MAY26-6L3ONX

Release Date
2026.05.20

PRODUCT INFORMATION

Product Name	CJC-1295 + Ipamorelin Blend
Dosage Form	Lyophilized powder (reconstitute before use)
Total Peptide Content	10 mg per vial (CJC-1295: 5 mg + Ipamorelin: 5 mg)
Appearance	White to off-white lyophilized cake/powder
Solubility	Freely soluble in bacteriostatic water or sterile water
Storage	Refrigerate 2-8 C; protect from light
Intended Use	Laboratory research use only - not for human consumption

COMPOSITION REFERENCE

CJC-1295 with DAC: Tyr-D-Ala-Asp-Ala-Ile-Phe-Thr-Gln-Ser-Tyr-Arg-Lys-Val-Leu-Ala-Gln-Leu-Ser-Ala-Arg-Lys-Leu-Leu-Gln-Asp-Ile-Leu-Ser-Arg-DAC

Ipamorelin: Aib-His-D-2-Nal-D-Phe-Lys-NH2

Peptide sequence records for identity verification purposes

QUALITY CONTROL TEST RESULTS

Test	Method	Specification	Result	Status
Appearance	Visual	White lyophilized cake	Conforms	PASS
Identity (CJC-1295)	HPLC/MS	Matches reference	Confirmed	PASS
Identity (Ipamorelin)	HPLC/MS	Matches reference	Confirmed	PASS
Purity (CJC-1295)	HPLC	>= 98.0%	99.1%	PASS
Purity (Ipamorelin)	HPLC	>= 98.0%	98.7%	PASS
Peptide Content	Gravimetric	Report result	12.04 mg	PASS
Endotoxin	USP <85> LAL	< 0.50 EU/mg	< 0.10 EU/mg	PASS
Sterility	USP <71>	No growth (14 days)	No growth	PASS

CERTIFICATION

This product has been manufactured, tested, and released in accordance with national quality standards. All tests performed meet or exceed applicable specifications.

RELEASED FOR DISTRIBUTION • LOT: CJCIPM-MAY26-6L3ONX • RELEASE DATE: 2026.05.20

Tirzepatide

QUALITY CONTROL CERTIFICATE

Tirzepatide (GIP/GLP-1 Dual Receptor Agonist)



Certificate No.
CP-QCC-2026-052639

Lot Number
TZ-MAY26-3V9ONX

Release Date
2026.05.27

PRODUCT INFORMATION

Product Name	Tirzepatide
Dosage Form	Lyophilized powder (reconstitute before use)
Peptide Content	10 mg per vial
Appearance	White to off-white lyophilized powder
Solubility	Freely soluble in sterile water or bacteriostatic water
Storage	Refrigerate 2-8 C; protect from light and moisture
Intended Use	Laboratory research use only - not for human consumption

COMPOSITION REFERENCE

Y-Aib-E-G-T-F-T-S-D-Y-S-I-Aib-L-D-K-I-A-Q-K(γE-AEEA-AEEA-C20diacid)-A-F-V-Q-W-L-I-A-G-G-P-S-S-G-A-P-P-P-S-NH2

39-mer GIP/GLP-1 dual agonist; C20 fatty diacid at K20 via γE-AEEA-AEEA linker; C-terminal amide; confirmed by LC-MS

QUALITY CONTROL TEST RESULTS

Test	Method	Specification	Result	Status
Appearance	Visual	White lyophilized powder	Conforms	PASS
Identity	HPLC/MS	Matches reference	Confirmed	PASS
Purity	HPLC	>= 98.0%	99.2%	PASS
Peptide Content	Gravimetric	Report result	10.1 mg	PASS
Related Substances	HPLC	<= 2.0%	0.8%	PASS
Endotoxin	USP <85> LAL	< 0.50 EU/mg	< 0.10 EU/mg	PASS
Sterility	USP <71>	No growth (14 days)	No growth	PASS

CERTIFICATION

This product has been manufactured, tested, and released in accordance with national quality standards. All tests performed meet or exceed applicable specifications.

RELEASED FOR DISTRIBUTION • LOT: TZ-MAY26-3V9ONX • RELEASE DATE: 2026.05.27