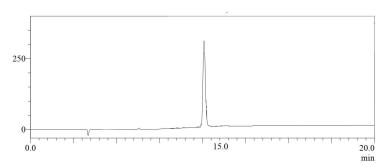


CERTIFICATE OF ANALYSIS

SAMPLE INFORMATION

Product Name	Tesamorelin 10 mg
Client Name/Lot No.	Glacier Aminos / Batch# TES1092501
Sequence	trans-hexenoyl-acid-Tyr-Ala-Asp-Ala-Ile-Phe-Thr-Asn-Ser-Tyr-Arg-Lys-Val-Leu-Gly-Gln-Leu-Ser-Ala-Arg-Lys-Leu-Leu-Gln-Asp-Ile-Met-Ser-Arg-Gln-Gln-Gly-Glu-Ser-Asn-Gln-Glu-Arg-Gly-Ala-Arg-Ala-Arg-Leu-NH2
Dissolution condition	100% H2O
Length	44AA
Molecular Weight	5135.86 g/mol

CHROMATOGRAM



Peak #	Ret. Time	Area %
1	14.903	99.640

TEST RESULTS

1E31 RESULTS				
	Specifications	Results		
Strength	10.00 mg	5.27 mg		
Appearance	White to off white lyophilized powder	Conforms		
Purity	≥98.0%	99.6%		
pH value	6.0-8.0	7.0		
Impurity	Single Impurity ≤1.0%	0.2%		
	Total Impurity ≤2.0%	0.4%		

TEST PARAMETERS

Pump A	0.1% trifluoroacetic in 100% water
Pump B	0.1% trifluoroacetic in 100% acetonitrile
Total Flow	1.0ml/min
Wavelength	214nm
Analytical Column	Agilent ZORBAX StableBond 5μm
Туре	C18 (4.6*250mm*5 µm)
Dissolution Method	100% H2O
Injection Volume	30uL

CONCLUSION

One 3ml vial contained a white lyophilized powder and has a purple cap with a silver crimp.

The sample was analysed using Reverse Phase High Performance Liquid Chromatography (RP-HPLC) and determined to contain 99.6% tesamorelin (5.27 mg), and the rest are impurities of minor significance.

CERTIFIED BY:

Dane Fredericksen Analytical Chemist 09/21/2025



