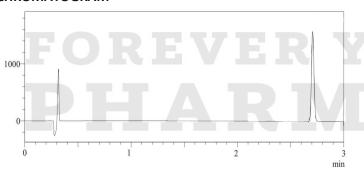


CERTIFICATE OF ANALYSIS

SAMPLE INFORMATION

Product Name	Semaglutide 5mg
Client Name/Lot No.	Encompass Peptides
Sequence	H-His-Aib-Glu-Gly-Thr-Phe-Thr-Ser-Asp-Val-Ser-Ser-Tyr-Leu-Glu-Gly-Gln-Ala-Ala-Lys(C18-diacid-gamma-Glu-OEG-OEG)-Glu-Phe-Ile-Ala-Trp-Leu-Val-Arg-Gly-Arg-Gly-OH
Dissolution condition	100% H2O
Length	31AA
Molecular Weight	4113.6 g/mol

CHROMATOGRAM



Peak #	Ret. Time	Area %
1	2.729	98.820

TEST RESULTS

	Specifications	Results
Strength	5.00 mg	5.59 mg
Appearance	White to off white lyophilized powder	Conforms
Purity	≥98.0%	98.8%
pH value	6.0-8.0	7.0
Impurity	Single Impurity ≤1.0%	0.9%
	Total Impurity ≤2.0%	1.2%

TEST PARAMETERS

Pump A	0.1% trifluoroacetic in 100% water			
Pump B	0.1% trifluoroacetic in 100% acetonitrile			
Total Flow	1.0ml/min			
Wavelength	220nm			
Analytical Column	Agilent ZORBAX StableBond 5μm			
Туре	C18 (4.6*250mm*5 µm)			
Dissolution Method	100% H2O			
Injection Volume	30uL			

CONCLUSION

One 3ml contained a white lyophilized powder and has a yellow cap and a silver crimp.

The sample was analysed using Reverse Phase High Performance Liquid Chromatography (RP-HPLC) and determined to contain 98.8% semaglutide (5.59 mg), and the rest are impurities of minor significance.

CERTIFIED BY:

Date D.

Dane Fredericksen Analytical Chemist 07/23/2025

