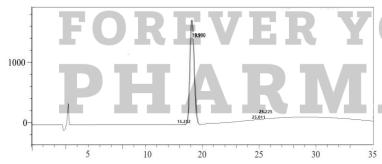


CERTIFICATE OF ANALYSIS

SAMPLE INFORMATION

Product Name	Retatrutide 10 mg	
Client Name/Lot No.	Forever Young Pharmacy / Lot# 224991	
Sequence	Tyr-{Aib}-Gln-Gly-Thr-Phe-Thr-Ser-Asp-Tyr-Ser-Ile-{α-Me-Leu}-Leu-Asp-Lys-{diacid-C20-gamma-Glu-(AEEA)-Lys}- Ala-Gln-{Aib}- Ala-Phe-Ile-Glu-Tyr-Leu-Leu-Glu-Gly-Gly-Pro-Ser-Ser-Gly-Ala- Pro-Pro-Pro-Ser-NH2 (sodium salt)	
Dissolution condition	1 100% H2O	
Length	39AA	
Molecular Weight	4813.6 g/mol	

CHROMATOGRAM



Г	Peak#	Ret. Time	Area %
	1	15.252	0.104
	2	19.990	99.611
	3	25.011	0 .115
	4	25.225	0.170

TEST RESULTS

	Specifications	Results
Strength	10.00 mg	10.55 mg
Appearance	White to off white crystallized 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	220nm			
Analytical Column Type	Agilent ZORBAX StableBond 5μm 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 yellow cap with silver foil and is Lot# 224991.

The sample was analysed using Reverse Phase High Performance Liquid Chromatography (RP-HPLC) and determined to contain 99.6% retatrutide (10.55 mg), and the rest are impurities of minor significance.

CERTIFIED BY:

Dajedo.

Dane Fredericksen Analytical Chemist