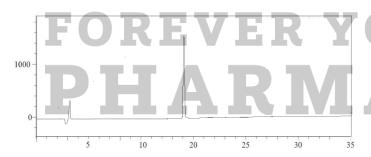


# **CERTIFICATE OF ANALYSIS**

## **SAMPLE INFORMATION**

Product Name	Retatrutide 30 mg	
Client Name/Lot No.	Forever Young Pharmacy / Lot# 25RET3001	
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	100% H2O	
Length	39AA	
Molecular Weight	4813.6 g/mol	

### **CHROMATOGRAM**



Peak#	Ret. Time	Area %
1	18.800	0.111
2	19.990	99.812
3	19.995	0.069
4	20.130	0.008

#### **TEST RESULTS**

	Specifications	Results
Strength	30.00 mg	31.05 mg
Appearance	White to off white crystallized powder	Conforms
Purity	≥98.0%	99.8%
pH value	6.0-8.0	8.0
Impurity	Single Impurity ≤1.0%	0.1%
	Total Impurity ≤2.0%	0.2%

### **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 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 clear cap with bronze crimp and is Lot# 25RET3001.

The sample was analysed using Reverse Phase High Performance Liquid Chromatography (RP-HPLC) and determined to contain 99.8% retatrutide (31.05 mg), and the rest are impurities of minor significance.

**CERTIFIED BY:** 

Dufando.

Dane Fredericksen Analytical Chemist

