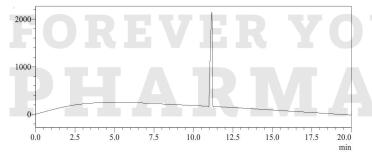


CERTIFICATE OF ANALYSIS

SAMPLE INFORMATION

r		
Product Name	LL-37 5mg	
Client Name/Lot No.	Forever Young Pharmacy / Lot# 25LL02	
Sequence	Leu-Leu-Gly-Asp-Phe-Phe-Arg-Lys-Ser-Lys-Glu-Lys-Ile-Gly-Lys-Glu-Phe-Lys-Arg-Ile-Val-Gln-Arg-Ile-Lys-Asp-Phe- Leu-Arg-Asn-Leu-Val-Pro-Arg-Thr-Glu-Ser	
Dissolution condition	100% H2O	
Length	37AA	
Molecular Weight	449.3 g/mol	

CHROMATOGRAM



Peak #	Ret. Time	Area %
1	5.674	0.149
2	10.932	0.141
3	11.292	99.546
4	11.607	0.164

TEST RESULTS

	Specifications	Results
Strength	5. 00 mg	5.70 mg
Appearance	White to off white lyophilized powder	Conforms
Purity	≥98.0%	99.5%
pH value	6.0-8.0	7.0
Impurity	Single Impurity ≤1.0%	0.2%
	Total Impurity ≤2.0%	0.5%

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 red flip off cap and silver crimp and Lot# 25LL02.

The sample was analysed using Reverse Phase High Performance Liquid Chromatography (RP-HPLC) and determined to contain 99.5% LL-37 (5.70 mg), and the rest are impurities of minor significance.

CERTIFIED BY:

Derfrencho

Dane Fredericksen Analytical Chemist



Verify the validity of test results by contacting support@foreveryoungpharmacy.com