

## CERTIFICATE OF ANALYSIS

### SAMPLE INFORMATION

<b>Product Name</b>	Tesamorelin 2 mg
<b>Sequence</b>	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-Gln-Gln-Gly-Ala-Glu-Gln-Gln-Gly-Ala-Gln-Glu-Gln-Gln-Gly-Ser-Gln-Glu-Gln-Gln-Gly-Glu-Glu-Gln-Gln-Gly-Glu-Glu-Gln-Gln-Gly-OH
<b>Dissolution Condition</b>	100% H <sub>2</sub> O
<b>Molecular Weight</b>	5135 g/mol

**CHROMATOGRAM:** Available on Request

**STORAGE CONDITION:** 0 °C

### PUMP SETTINGS

<b>Pump A</b>	HPLC Grade water with 0.1% Formic Acid
<b>Pump D</b>	100% Acetonitrile with 0.1% Formic Acid
<b>Column Usage</b>	Agilent Zorbax 300SB-C18, C18, 2.1 × 100 mm, 1.7 µm, 300 Å
<b>Gradient</b>	25–60% ACN over 20 minutes
<b>Pump Settings</b>	0.5 ml/min
<b>Injection Volume:</b>	10 µl
<b>Temperature</b>	35 °C

### PURITY RESULT

Purity is **99.6%**

Total impurity 0.4% of inert solvents; no significant related substances or degradation products detected above 0.1%.

### CONCLUSION

One 3 ml vial contained a white lyophilized powder with a silver crimp and is labeled batch **#27231**.

Sample strength: **2 mg**

Sample weight in grams: **1.9**

This sample was analyzed on a Thermo Ulitimate U3000 HPLC stack using reverse phased high performance liquid chromatography and determined to contain **99.6% Tesamorelin 2 mg**, with the rest being impurities of minor significance

### CERTIFICATION

**Certified by:** Jonathan Barber

**Title:** Analytical Chemist

**Date:** 23 July 2025

