ASCENDTIALS LLC
Research compounds · Quality Control Department

CERTIFICATE OF ANALYSIS

Document COA-KLW-2501C

KLOW Blend

LOT NUMBER
KLW-2501C

DATE OF ANALYSIS
August 21, 2026

RETEST DATE
August 20, 2028

ANALYTICAL LABORATORY
Third-party HPLC/MS

ANALYTICAL RESULTS

TEST	METHOD	SPECIFICATION	RESULT
Appearance	Visual	White to off-white lyophilized powder	**Conforms**
Identity (molecular mass)	ESI-MS	Conforms to structure	**Conforms**
Purity	RP-HPLC, UV 214 nm	NLT 98.0%	**98.50%**
Single largest impurity	RP-HPLC, UV 214 nm	NMT 1.0%	**0.58%**
Water content	Karl Fischer titration	NMT 8.0%	**4.2%**
Acetate content	Ion chromatography	NMT 15.0%	**9.1%**
Peptide content	Nitrogen determination	NLT 80.0%	**89.5%**
Residual solvents	GC headspace, ICH Q3C	Meets ICH limits	**Conforms**
Bacterial endotoxins	LAL, USP <85>	< 1.0 EU/mg	**Conforms**

CONFORMS — lot KLW-2501C meets all specifications listed above.

CHROMATOGRAPHIC CONDITIONS

Column: C18 reversed phase, 4.6 × 250 mm, 5 µm, 100 Å
Mobile phase A: 0.1% TFA in water · B: 0.1% TFA in acetonitrile
Gradient: 5-65% B over 30 min · Flow: 1.0 mL/min · 25 °C
Detection: UV 214 nm · Injection volume: 20 µL
Mass spectrometry: ESI, positive ion mode

STORAGE & HANDLING

Lyophilized: 2-8°C, protected from light.

RELEASED BY
Quality Control, Ascendtials LLC

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This certificate applies only to the lot identified above. Not for use in food, drugs, medical devices, or cosmetics.