

CERTIFICATE OF ANALYSIS / BATCH RELEASE & IDENTITY VERIFICATION

KPV 10mg

Lys-Pro-Val synthetic tripeptide (KPV) · sterile-filled lyophilised powder, released against independent third-party HPLC, LC-MS and bacterial-endotoxin analysis.

PRODUCT KPV 10mg Lyophilised powder	LOT · BATCH PH-kv101123 Single production lot	TESTING LABORATORY Independent Third-party laboratory	REPORT DATE 20 April 2026
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HPLC-UV PURITY 99.74% Mean of two vials · 99.74% / 99.73% ● MEETS RELEASE SPEC	IDENTITY LC-MS confirmation	KPV Confirmed
	NET PEPTIDE CONTENT Mean of two vials	11.34mg ≥ 10mg
	BACTERIAL ENDOTOXIN LAL · USP <85>	Pass ≤0.05 EU/mL

01 PURITY · IDENTITY · CONTENT · ENDOTOXIN

Analytical Results

TEST	METHOD	RESULT	RELEASE SPECIFICATION	STATUS
Identity Peptide confirmation	LC-MS	KPV · [M+H] ⁺ m/z 343.2	Conforms to reference mass	PASS
Chromatographic Purity Main peak, area %	HPLC-UV	99.74% Vial 1 · 99.74% Vial 2 · 99.73%	≥ 98.0%	PASS
Net Peptide Content Mean of two vials	Gravimetric	11.34mg Vial 1 · 11.40mg Vial 2 · 11.28mg	≥ 10.0mg (label claim)	PASS
Bacterial Endotoxin Replicate 1 & 2	LAL · USP <85>	Pass / Pass Assay sensitivity ≤ 0.05 EU/mL	≤ 0.05 EU/mL	PASS
Appearance Visual inspection	Visual	White lyophilised powder Sterile-filled, sealed vial	White cake / powder	PASS

Methods. Purity by high-performance liquid chromatography with UV detection (HPLC-UV); identity by liquid chromatography-mass spectrometry (LC-MS). Bacterial endotoxin by Limulus Amebocyte Lysate (LAL) assay in accordance with USP <85> under validated laboratory conditions, assay sensitivity ≤ 0.05 EU/mL. **Status** is assessed against acceptance criteria set by Eira London for research-grade (RUO) release.

02 INSTRUMENT TRACES

Chromatogram & Mass Confirmation

CHROMATOGRAM	HPLC-UV / PDA	MASS CONFIRMATION	LC-MS
	Single dominant peak eluting at ~1 min, integrated at 99.74% of total area. No significant co-eluting impurities.		Protonated molecular ion [M+H] ⁺ observed at m/z 343.2 (with [M+2H] ²⁺ at m/z 172.1), consistent with the KPV reference standard.

Released for research use. This batch conforms to Eira London's research-grade release specification on the basis of the independent analytical data above. Certificate refers only to the lot and sample(s) tested.

Eira London

AUTHORISED — QUALITY & COMPLIANCE
Batch released 20 April 2026

