Certificate of Analysis

ILS Laboratories
8222 Vickers St, Suite 106, San Diego, CA 92111
(619) 329-3999 | ils-lab.com

TESA/IPA - 13mg

PASS

Tested for: PeptidePharm
PeptidePharm.com

COA #:	**COA-2026-412647**	Method:	**Full QC Panel**
Lot Number:	**TP13-05042026**	Analysis Date:	**05/08/2026**
Accession #:	**ACC-2026-2046**	Appearance:	**Good**
Labeled Content:	**13mg**	Sample Matrix:	**Lyophilized**
		Date Received:	**05/05/2026**

Scan to verify authenticity at ils-lab.com
Access Code: XZDYZC36

Identity	Peptide Purity
TESA/IPA	**99.82%**

TESA/IPA 13mg - TP13-05042026

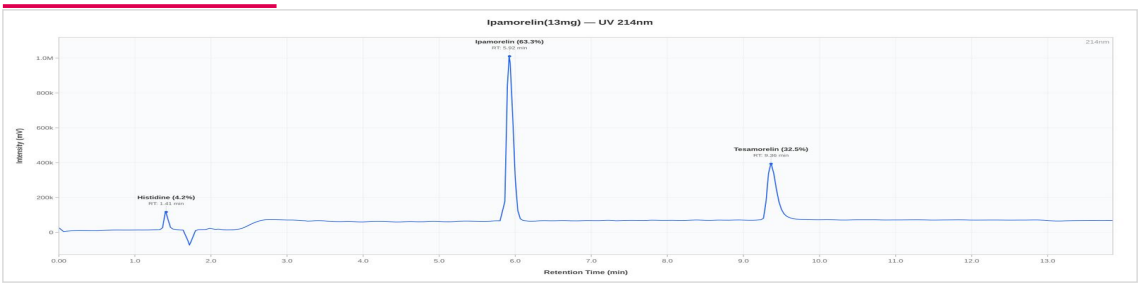
Blend Purity & Quant (HPLC)

Analyte	Specification	Result	Unit	Status
Peptide Purity (HPLC)	>= 95.0%	**99.82%**	%	**PASS**
Identity (HPLC-RTM)	TESAMORELIN + IPAMORELIN	**Confirmed**	-	**PASS**

Additional Tests

Analyte	Specification	Result	Unit	Status
Net Blend Peptide Content	Report Only	**13.74**	mg	N/A

HPLC Chromatogram



Representative chromatogram, Dedicated V0 (99.82% purity, closest to batch mean of 99.81%)

HPLC Conformity Testing Results (2 samples tested)

Sample	Purity	NPC	ID	Result
Dedicated V0	99.82%	13.74 mg	Confirmed	**PASS**
Conformity V1	99.80%	13.55 mg	Confirmed	**PASS**
Mean	**99.81%**	**13.64 mg**	—	—
Std Dev	**0.0100%**	**0.0950 mg**	—	—

Dr. Greg Kalyuzhny
Lab Director
6/2/2026

COA #: **COA-2026-412647**
Access Code: **XZDYZC36**
Verify: portal.ils-lab.com/verify/cDpLYikJ-SASsG7_
Issued: 6/2/2026

Heavy Metals Analysis (ICP-MS)

Test	Specification	Result	Status
Arsenic (As)	NMT 1.5 ppm	*Not Detected*	**PASS**
Cadmium (Cd)	NMT 0.5 ppm	*Not Detected*	**PASS**
Chromium (Cr)	NMT 10 ppm	*Not Detected*	**PASS**
Mercury (Hg)	NMT 1.5 ppm	*Not Detected*	**PASS**
Lead (Pb)	NMT 1 ppm	*Not Detected*	**PASS**

Sterility Testing (PCR)

Test	Specification	Result	Status
Sterility (PCR)	No Growth	**No Growth**	**PASS**

Endotoxin Testing (USP <85>)

Test	Specification	Result	Status
Endotoxin (USP <85>)	Report Result	**0.085 EU/mL**	**Reported**

About this result: Endotoxin is reported as a quantitative value. Acceptable limits vary by product type and matrix, so no universal pass/fail threshold applies to RUO products. This result is below commonly referenced endotoxin thresholds.

Notes & Methodology

1. Date Tested: 06/02/2026. Methods: Blend Purity & Quant (HPLC); Additional Tests.
2. The sample was confirmed to be TESA/IPA by HPLC. Identification by chromatographic retention time comparison with a reference standard.
3. Elemental impurities analyzed by ICP-MS per USP <233> methodology. Acceptance criteria are internal laboratory quality screening limits for research-use materials and do not represent evaluation against any specific pharmacopeial monograph or product specification.
4. Endotoxin tested per USP <85> kinetic turbidimetric method. Acceptance criteria per client specification.
5. Peptide purity determined by RP-HPLC area normalization at 214 nm. Value represents the percentage of the target peptide relative to all peptide-related peaks. Non-peptide process-related impurities, if detected, are excluded from the calculation.
6. Chromatogram shown is representative: Dedicated V0 (99.82% purity, closest to batch mean of 99.81%).

ilac-MRA PJLA Testing Accreditation No. 59423

Dr. Greg Kalyuzhny
Lab Director
6/2/2026

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