Certificate of Analysis

ILS Laboratories
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(619) 329-3999 | ils-lab.com

Tesamorelin / Ipamorelin - 20

Tested for: px1 research
PX1RESEARCH.COM

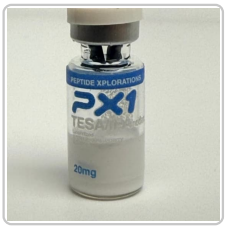
PASS

Scan to verify authenticity at ils-lab.com
Access Code: RZ2JTR3V

COA #:	**COA-2026-ANSK9X**	Analysis Date:	**10/07/2026**
Lot Number:	**PX1TESAIPA20-0021**	Appearance:	**Good**
Labeled Content:	**20**	Volume:	**3**
Volume:	**3**	Date Received:	**09/30/2026**

Method: **Full QC Panel**

Identity	Peptide Purity	
Tesamorelin / Ipamorelin	**99.90%**	Fentanyl Free



Tesamorelin / Ipamorelin 20 - PX1TESAIPA20-0021

Full QC Panel

Analyte	Specification	Result	Unit	Status
Peptide Purity (HPLC)	>= 95.0%	**99.90%**	%	**PASS**
Net Blend Peptide Content	Report Only	**21.21**	mg	N/A
-- TESAMORELIN		*10.51*	*mg*	
-- IPAMORELIN		*10.70*	*mg*	
Identity (HPLC-RTM)	TESAMORELIN + IPAMORELIN	**Confirmed**	-	**PASS**
Fentanyl Screen	Immunoassay, 50 ng/mL cutoff	**Not Detected**	-	**PASS**

HPLC Chromatogram



Tesamorelin / Ipamorelin 20 - PX1TESAIPA20-0021: UV Chromatogram

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>)	NMT 0.25 EU/mL	**0.189 EU/mL**	**PASS**
Acceptance criteria: NMT 0.25 EU/mL per client specification.			

Notes & Methodology

1. Date Tested: 10/07/2026. Methods: Full QC Panel.
2. The sample was confirmed to be Tesamorelin / Ipamorelin 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 chromogenic 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. Purity values are capped at 99.9% to reflect instrument limit of detection (LOD). Trace impurities below the detection threshold may be present.
7. Per-component content determined by individual peak integration. Not all labeled components may be present.
8. Accreditation Scope: ILS Laboratories maintains ISO/IEC 17025 accreditation for activities listed in its published PJLA scope. Peptide testing reported on this certificate is performed for Research Use Only.
9. This certificate reports analytical results for the sample(s) submitted to ILS Laboratories for testing. All sample identifications, product names, brand names, lot/batch numbers, and labeled content are provided by the submitting party and have not been independently verified. Results reflect only the analytical testing of the specific sample(s) provided and do not constitute an evaluation or endorsement of the manufacturer's processes, quality systems, or overall production. ILS Laboratories makes no representations regarding the intended use, safety, efficacy, regulatory status, or intellectual property ownership of any materials tested.

Issued by ILS Laboratories
(619) 329-3999
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Lab Director
10/07/2026

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