Certificate of Analysis

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

Tesamorelin - 20mg

PASS

Tested for: Crush Research
https://www.crushresearch.shop/

COA #:	**COA-2026-YSMFIF**	Method:	**Full QC Panel**
Lot Number:	**CR-TSM20-0307-1**	Analysis Date:	**05/11/2026**
Accession #:	**ACC-2026-2073**	Appearance:	**Good**
Concentration:	**20mg**	Volume:	**3mL**
Sample Matrix:	**Powder**	Received:	**05/06/2026**

Scan to verify authenticity at ils-lab.com

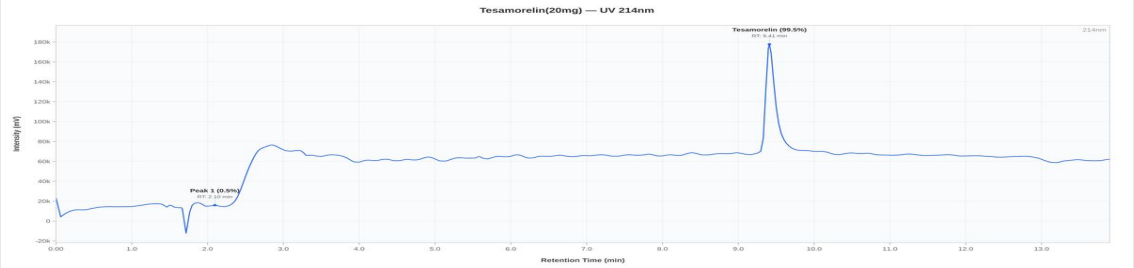
Identity	Purity
Tesamorelin	**99.47%**

Tesamorelin 20mg - CR-TSM20-0307-1

Purity & Quant (HPLC)

Analyte	Specification	Result	Unit	Status
Purity (HPLC)	>= 95.0%	**99.47%**	%	**PASS**
Net Peptide Content	Report Only	**21.32**	mg	N/A
Identity (ID)	Tesamorelin	**Confirmed**	-	**PASS**

HPLC Chromatogram



Representative chromatogram, Dedicated V0 (99.46% purity, closest to batch mean of 99.37%)

HPLC Conformity Testing Results (3 samples tested)

Sample	Purity	NPC	ID	Result
Dedicated V0	99.46%	20.67 mg	Confirmed	**PASS**
Conformity V1	99.17%	20.8 mg	Confirmed	**PASS**
Conformity V2	99.47%	21.32 mg	Confirmed	**PASS**
Mean	**99.37%**	**20.93 mg**	—	—
Std Dev	**0.1391%**	**0.2808 mg**	—	—

Dr. Greg Kalyuzhny
Lab Director
5/11/2026

COA #: **COA-2026-YSMFIF**
Access Code: **O4LWWOSB**
Verify: portal.ils-lab.com/verify/uiwjOpEU_Lt18Scq
Issued: 5/11/2026

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Sample Matrix:	**Powder**	Received:	**05/06/2026**

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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	Unit	Status
Sterility (PCR)	No Growth	**No Growth**	-	**PASS**

Endotoxin Testing (USP <85>)

Test	Specification	Result	Unit	Status
Endotoxin (USP <85>)	< 0.25 EU/mL	**0.09 EU/mL**		**PASS**

Notes & Methodology

1. Date Tested: 05/11/2026. Methods: Purity & Quant (HPLC).
2. The sample was confirmed to be Tesamorelin 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 route-of-administration standard.
4. Chromatogram shown is representative: Dedicated V0 (99.46% purity, closest to batch mean of 99.37%).

Dr. Greg Kalyuzhny
Lab Director
5/11/2026

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Verify: portal.ils-lab.com/verify/uiwjOpEU_Lt18Scq
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