Certificate of Analysis

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

Triple Agonist - 50mg

PASS

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

COA #:	**COA-2026-I8BEKL**	Method:	**Purity & Quant (HPLC)**
Lot Number:	**CR26043XAG50-01**	Analysis Date:	**05/27/2026**
Accession #:	**ACC-2026-2999**	Appearance:	**Good**
Concentration:	**50mg**	Volume:	**3mL**
Sample Matrix:	**Lyophilized**	Received:	**05/22/2026**

Scan to verify authenticity at ils-lab.com

Identity	Purity
Triple Agonist	**99.37%**

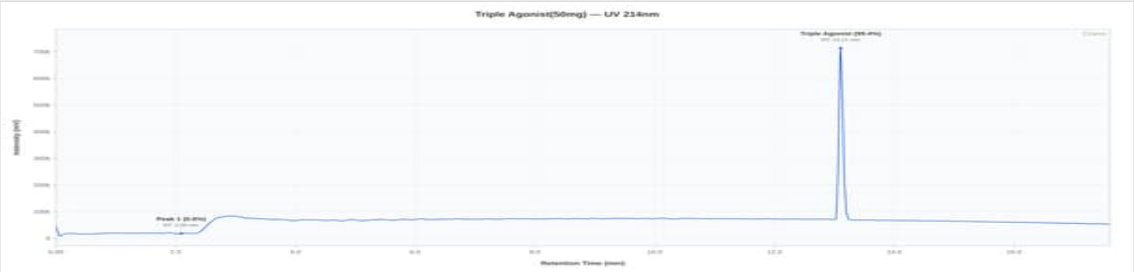


Triple Agonist 50mg - CR26043XAG50-01

Purity & Quant (HPLC)

Analyte	Specification	Result	Unit	Status
Purity (HPLC)	>= 95.0%	**99.37%**	%	**PASS**
Net Peptide Content	Report Only	**53.27**	mg	N/A
Identity (ID)	Triple Agonist	**Confirmed**	-	**PASS**

HPLC Chromatogram



Representative chromatogram, Conformity V2 (99.36% purity, closest to batch mean of 99.31%)

HPLC Conformity Testing Results (3 samples tested)

Sample	Purity	NPC	ID	Result
Dedicated V0	99.37%	53.27 mg	Confirmed	**PASS**
Conformity V1	99.21%	51.48 mg	Confirmed	**PASS**
Conformity V2	99.36%	51.24 mg	Confirmed	**PASS**
Mean	**99.31%**	**52.00 mg**	—	—
Std Dev	**0.0732%**	**0.9057 mg**	—	—

Dr. Greg Kalyuzhny
Lab Director
5/27/2026

COA #: **COA-2026-I8BEKL**
Access Code: **KDO91Y6-**
Verify: portal.ils-lab.com/verify/2txyH3tlQ2PRs7Ro
Issued: 5/27/2026

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Lot Number:	**CR26043XAG50-01**	Analysis Date:	**05/27/2026**
Accession #:	**ACC-2026-2999**	Appearance:	**Good**
Concentration:	**50mg**	Volume:	**3mL**
Sample Matrix:	**Lyophilized**	Received:	**05/22/2026**

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

Endotoxin Testing (USP <85>)

Test	Specification	Result	Status
Endotoxin (USP <85>)	Report Result	**NMT 0.05 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: 05/27/2026. Methods: Purity & Quant (HPLC).
2. The sample was confirmed to be Triple Agonist 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. Chromatogram shown is representative: Conformity V2 (99.36% purity, closest to batch mean of 99.31%).

ilac-MRA PJLA Testing Accreditation No. 59423

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