Certificate of Analysis

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

Double Agonist - 20mg

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

PASS

COA #:	**COA-2026-JCGGMS**	Method:	**Full QC Panel**
Lot Number:	**CR-2XAG20-2605-1**	Analysis Date:	**06/01/2026**
Accession #:	**ACC-2026-3292**	Appearance:	**Good**
Labeled Content:	**20mg**	Sample Matrix:	**Lyophilized**
		Date Received:	**05/27/2026**

Scan to verify authenticity at ils-lab.com
Access Code: YFUPGRB6

Identity	Peptide Purity	
Double Agonist	**99.39%**	Fentanyl Free

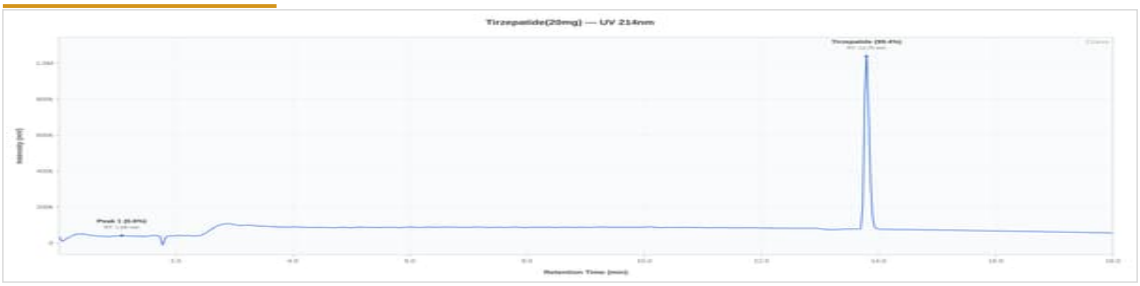


Double Agonist 20mg - CR-2XAG20-2605-1

Purity & Quant (HPLC)

Analyte	Specification	Result	Unit	Status
Peptide Purity (HPLC)	>= 95.0%	**99.39%**	%	**PASS**
Net Peptide Content	Report Only	**20.91**	mg	N/A
Identity (HPLC-RTM)	Double Agonist	**Confirmed**	-	**PASS**
Fentanyl Screen	Immunoassay, 50 ng/mL cutoff	**Not Detected**	-	**PASS**

HPLC Chromatogram



Representative chromatogram, Dedicated V0 (99.39% purity, closest to batch mean of 99.30%)

HPLC Conformity Testing Results (2 samples tested)

Sample	Purity	NPC	ID	Result
Dedicated V0	99.39%	20.91 mg	Confirmed	**PASS**
Conformity V1	99.21%	20.61 mg	Confirmed	**PASS**
Mean	**99.30%**	**20.76 mg**	—	—
Std Dev	**0.0900%**	**0.1500 mg**	—	—

Dr. Greg Kalyuzhny
Lab Director
6/1/2026

COA #: **COA-2026-JCGGMS**
Access Code: **YFUPGRB6**
Verify: portal.ils-lab.com/verify/6dIiIViOcJGg9Z4n
Issued: 6/1/2026

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>)	NMT 5 EU/mL	**NMT 0.05 EU/mL**	**PASS**

Acceptance criteria: NMT 5 EU/mL per client specification.

Notes & Methodology

1. Date Tested: 06/01/2026. Methods: Purity & Quant (HPLC).
2. The sample was confirmed to be Double 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. 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 (99.39% purity, closest to batch mean of 99.30%).

Dr. Greg Kalyuzhny
Lab Director
6/1/2026

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