

## ILS Laboratories

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

## CJC-1295 No DAC (10mg) - 10mg

Tested for: Kovagen

www.Kovagen.com

COA #: COA-2026-954801  
Lot Number: IE701096  
Accession #: ACC-2026-13938  
Labeled Content: 10mg

Analysis Date: 08/03/2026  
Appearance: Good  
Sample Matrix: Powder  
Volume: 3mL  
Date Received: 2026-08-03

**PASS**



Scan to verify  
authenticity at ils-lab.com  
Access Code: ZZHXGKJ6

Method: Full QC Panel

Identity

**CJC-1295 No DAC (10mg)**

Peptide Purity

**99.79%**



Fentanyl  
Free

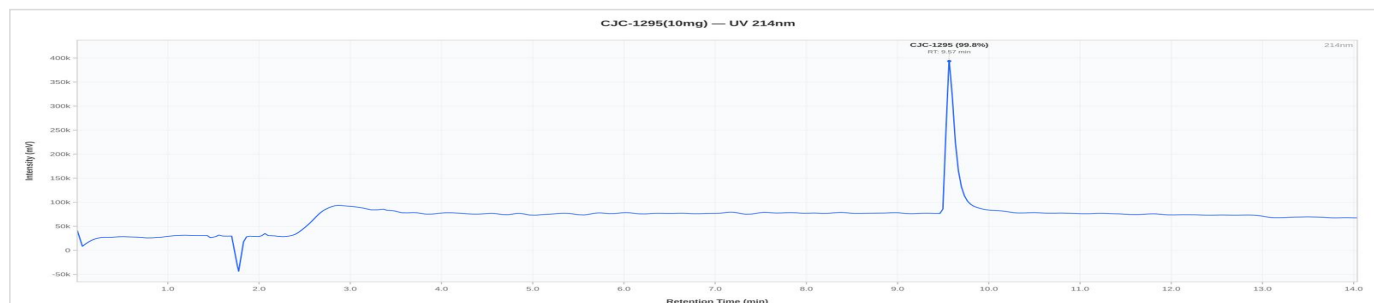
## Purity, Identity & Quantitation (HPLC)

| Analyte                   | Specification                | Result       | Unit | Status |
|---------------------------|------------------------------|--------------|------|--------|
| Peptide Purity (HPLC)     | >= 95.0%                     | 99.79%       | %    | PASS   |
| Net Blend Peptide Content | Report Only                  | 10.27        | mg   | N/A    |
| -- CJC-1295               |                              | 5.13         | mg   |        |
| -- Ipamorelin             |                              | 5.13         | mg   |        |
| Identity (HPLC-RTM)       | CJC-1295 + IPAMORELIN        | Confirmed    | -    | PASS   |
| Fentanyl Screen           | Immunoassay, 50 ng/mL cutoff | Not Detected | -    | PASS   |



CJC-1295 No DAC (10mg) 10mg - IE701096

## HPLC Chromatogram



Representative chromatogram, Conformity V2



*Dr. Greg Kalyuzhny*

**Dr. Greg Kalyuzhny**  
Lab Director  
8/3/2026

COA #: COA-2026-954801  
Access Code: ZZHXGKJ6  
Verify: [portal.ils-lab.com/verify/e31GxS1UYICOsLVy](https://portal.ils-lab.com/verify/e31GxS1UYICOsLVy)  
Issued: 8/3/2026

HPLC Conformity Testing Results (6 samples tested)

| Sample        | Purity  | NPC       | ID        | Result |
|---------------|---------|-----------|-----------|--------|
| Dedicated V0  | 99.79%  | 10.27 mg  | Confirmed | PASS   |
| Dedicated V0  | 99.79%  | 10.27 mg  | Confirmed | PASS   |
| Conformity V1 | 99.16%  | 10.45 mg  | Confirmed | PASS   |
| Conformity V1 | 99.16%  | 10.45 mg  | Confirmed | PASS   |
| Conformity V2 | 99.78%  | 10.57 mg  | Confirmed | PASS   |
| Conformity V2 | 99.78%  | 10.57 mg  | Confirmed | PASS   |
| Mean          | 99.58%  | 10.43 mg  | —         | —      |
| Std Dev       | 0.2947% | 0.1233 mg | —         | —      |

Heavy Metals Analysis (ICP-MS)

| Test          | Specification | Result     | Status |
|---------------|---------------|------------|--------|
| Arsenic (As)  | NMT 1.5 ppm   | 0.0953 ppm | PASS   |
| Cadmium (Cd)  | NMT 0.5 ppm   | 0.0534 ppm | PASS   |
| Lead (Pb)     | NMT 1.0 ppm   | 0.1874 ppm | PASS   |
| Mercury (Hg)  | NMT 1.5 ppm   | 0.2015 ppm | PASS   |
| Chromium (Cr) | NMT 10.0 ppm  | 0.692 ppm  | 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 | 0.155 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: 08/03/2026. Methods: Purity, Identity & Quantitation (HPLC).
- 2. The sample was confirmed to be CJC-1295 No DAC (10mg) 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. Per-component content calculated from total net peptide content using the manufacturer's stated formulation ratio (CJC-1295, Ipamorelin). All components confirmed present by HPLC identity testing, unless explicitly stated otherwise.
- 7. Chromatogram shown is representative: Conformity V2 (99.78% purity, closest to batch mean of 99.58%).



  
Dr. Greg Kalyuzhny  
Lab Director  
8/3/2026

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