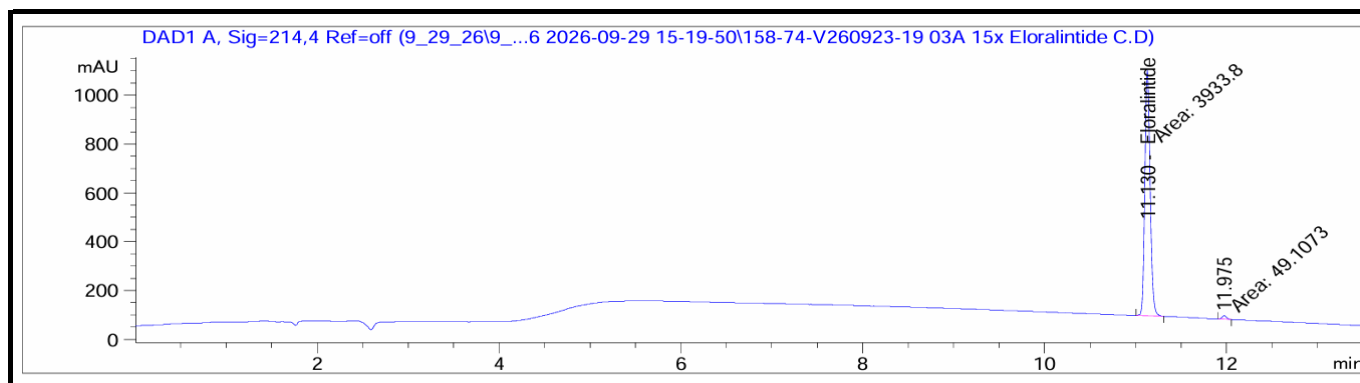


# Certificate of Analysis

Eloralintide 10 mg

Report To:  
**Chengdu Biotech**

Compound: Eloralintide  
Amount: 10 mg  
Laboratory ID: V260923-19 003  
Lot Number: CN-ELORA10-6SEP26-WHITE  
Date Released: 10/6/2026



| Analysis               | Method      | Result                       |          |          |
|------------------------|-------------|------------------------------|----------|----------|
| Identity               | HPLC-UV/VIS | Pass - Matches Ref. Standard |          |          |
| Chromatographic Purity | HPLC-UV/VIS | 98.77%                       | 98.75%   | 98.76%   |
| Quantity               | HPLC-UV/VIS | 17.84 mg                     | 15.42 mg | 15.84 mg |
| Fentanyl Screening     | Immunoassay | <50ng/mL                     |          |          |




Report By: Dustin Newman, Laboratory Director on 10/6/2026



Approved By: Tori Johnson, Operations Manager on 10/6/2026

Revised by HG 10/6/2026

Please consult A2LA Certificate #6377.01.01 for a list of accredited tests. Samples were received in acceptable condition. The result(s) in this report relate only to the portion of the sample(s) tested. All analyses were performed consistent with the Vanguard Laboratory Quality Management System. Vanguard Laboratory and its staff did not observe or participate in the sample selection process, and cannot confirm the authenticity of the sample or its representativeness of the associated lot/batch. Blend purity is the combined chromatographic area % of peaks identified as the intended components, not absolute mass purity. It does not establish individual component purity, quantity, or ratio. Co-eluting or otherwise undetected impurities may not be included, despite peak-purity and spectral-library review.



Vanguard Laboratory  
2635 Parkmont Ln  
Olympia, Wa 98502  
360-967-7010

# Certificate of Analysis

Eloralintide 10 mg

Report To:  
**Chengdu Biotech**

Compound: Eloralintide  
Amount: 10 mg  
Laboratory ID: V260923-19 003  
Lot Number: CN-ELORA10-6SEP26-WHITE  
Date Released: 10/6/2026

| Analyte      | Method | LOQ (ppm) | Result (ppm) |
|--------------|--------|-----------|--------------|
| Arsenic (As) | ICP-MS | 0.01      | ND           |
| Cadmium (Cd) | ICP-MS | 0.01      | ND           |
| Lead (Pb)    | ICP-MS | 0.01      | ND           |
| Mercury (Hg) | ICP-MS | 0.005     | ND           |

Run ID:

| Analysis   | Method   | Result             |
|------------|----------|--------------------|
| Endotoxins | LAL      | <5.00 EU/Vial      |
| Sterility  | USP <71> | No Growth Detected |

Run ID:

| Analysis                    | Method                | Result |
|-----------------------------|-----------------------|--------|
| Container Closure Integrity | Dye Ingress Method    | Pass   |
| Solubility Analysis         | Fully Soluble in 2 mL | Pass   |

Report By: Dustin Newman, Laboratory Director on 10/6/2026

Approved By: Tori Johnson, Operations Manager on 10/6/2026

ND: Non-Detect

LOD: Limit of Quantification



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