Certificate of Analysis

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

DSIP - 30mg

Tested for: Glow Aminos
www.glowaminos.com

PASS

COA #:	**COA-2026-412971**	Analysis Date:	**06/17/2026**
Lot Number:	**GL-010166**	Appearance:	**Good**
Accession #:	**ACC-2026-4797**	Sample Matrix:	**Liquid/Solution**
Labeled Content:	**30mg**	Vial Size:	**30mL**
		Date Received:	**06/12/2026**

Scan to verify authenticity at ils-lab.com
Access Code: JS8ULR8B

Method: **Full QC Panel**

Identity	Peptide Purity	
DSIP	**98.64%**	**Fentanyl Free**

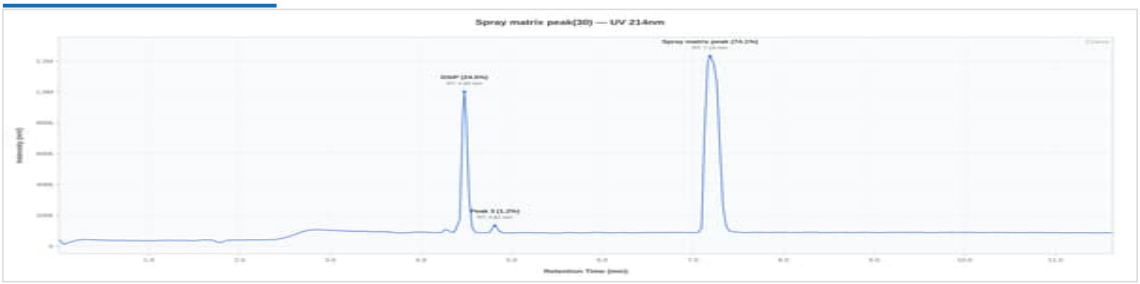


DSIP 30mg - GL-010166

Purity & Quant (HPLC)

Analyte	Specification	Result	Unit	Status
Peptide Purity (HPLC)	>= 95.0%	**98.64%**	%	**PASS**
Net Peptide Content	Report Only	**31.23**	mg	N/A
Identity (HPLC-RTM)	DSIP;Spray matrix peak	**Confirmed**	-	**PASS**
Fentanyl Screen	Immunoassay, 50 ng/mL cutoff	**Not Detected**	-	**PASS**

HPLC Chromatogram



Representative chromatogram, Conformity V1 (97.71% purity, closest to batch mean of 98.01%)

HPLC Conformity Testing Results (3 samples tested)

Sample	Purity	NPC	ID	Result
Dedicated V0	98.64%	31.23 mg	Confirmed	**PASS**
Conformity V1	97.71%	31.07 mg	Confirmed	**PASS**
Conformity V2	97.69%	31.31 mg	Confirmed	**PASS**
Mean	**98.01%**	**31.20 mg**	—	—
Std Dev	**0.4432%**	**0.0098 mg**	—	—

Dr. Greg Kalyuzhny
Lab Director
6/19/2026

COA #: **COA-2026-412971**
Access Code: **JS8ULR8B**
Verify: portal.ils-lab.com/verify/awOR6OF3JI6k0HPL
Issued: 6/19/2026

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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	**0.116 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: 06/19/2026. Methods: Purity & Quant (HPLC).
2. The sample was confirmed to be DSIP 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: Conformity V1 (97.71% purity, closest to batch mean of 98.01%).

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