

ILS Laboratories

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(619) 329-3999 | ils-lab.com

Semax/Selank/Pinealon - 60mg



Tested for: NextGen Peptides
www.NGpeptide.com

COA #: COA-2026-XQOBBU
Lot Number: NSP-SSP-01
Accession #: ACC-2026-7211
Labeled Content: 60mg

Analysis Date: 07/08/2026
Appearance: Good
Sample Matrix: Liquid/Solution
Date Received: 06/30/2026

PASS



Scan to verify
authenticity at ils-lab.com
Access Code: ZQJRQ52N

Method: Full QC Panel

Identity

Semax/Selank/Pinealon

Peptide Purity

99.63%

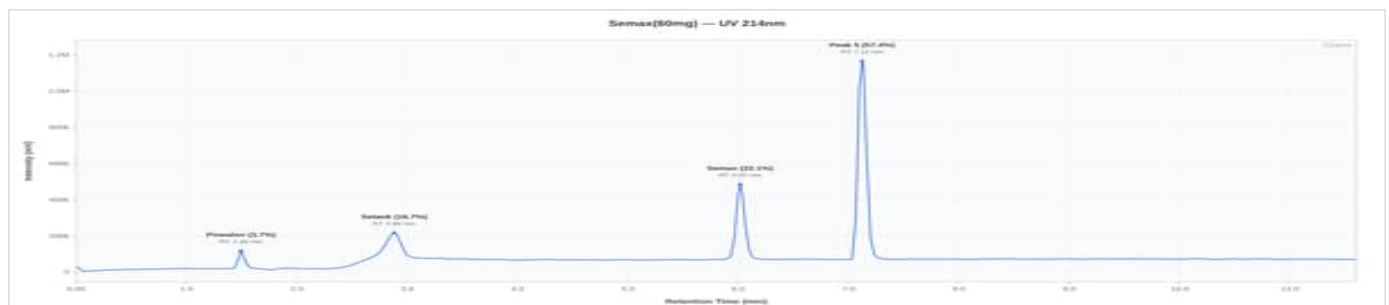


Fentanyl
Free

Full QC Panel

Analyte	Specification	Result	Unit	Status
Peptide Purity (HPLC)	>= 95.0%	99.63%	%	PASS
Net Blend Peptide Content	Report Only	63.59	mg	N/A
-- Semax		21.30	mg	
-- Selank		21.11	mg	
-- Pinealon		21.18	mg	
Identity (HPLC-RTM)	SEMAX + SELANK + PINEALON	Confirmed	-	PASS
Heavy Metals				
Arsenic (As)	<= 1.5 ppm	Not Detected	ppm	PASS
Cadmium (Cd)	<= 0.5 ppm	Not Detected	ppm	PASS
Lead (Pb)	<= 1.0 ppm	Not Detected	ppm	PASS
Mercury (Hg)	<= 1.5 ppm	Not Detected	ppm	PASS
Chromium (Cr)	<= 10.0 ppm	Not Detected	ppm	PASS
Fentanyl Screen	Immunoassay, 50 ng/mL cutoff	Not Detected	-	PASS

HPLC Chromatogram



Semax/Selank/Pinealon 60mg - NSP-SSP-01: UV Chromatogram



Dr. Greg Kalyuzhny

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Lab Director
7/8/2026

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Verify: portal.ils-lab.com/verify/qIRxsqCHXF25uJK0

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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	0.16 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: 07/08/2026. Methods: Full QC Panel.
2. The sample was confirmed to be Semax/Selank/Pinealon by HPLC. Identification by chromatographic retention time comparison with a reference standard.
3. Endotoxin tested per USP <85> kinetic turbidimetric method. Acceptance criteria per client specification.
4. 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.
5. Per-component content calculated from total net peptide content using the manufacturer's stated formulation ratio (Semax, Selank, Pinealon). All components confirmed present by HPLC identity testing, unless explicitly stated otherwise.




Dr. Greg Kalyuzhny
Lab Director
7/8/2026

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