

Customer:

EA Sample ID: 26EA0727-035

Date Received:

07/27/2026

Sample Name: Acetyl Selank

Date Completed:

Sample Type: Powder

08/04/2026

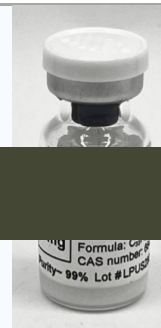
Batch/Lot: LPUS28012

ETHOS
ANALYTICS

CERTIFICATE OF ANALYSIS

Summary of Results

Analysis Type	Method	Date Tested	Status
N-Acetyl Selank	HPLC	08/04/2026	Complete
Bacterial Endotoxins	USP <85>	08/04/2026	Complete
Rapid Sterility	qPCR	08/04/2026	Complete
Heavy Metals	ICP-MS	07/30/2026	Pass
Microbiological	Petrifilm/qPCR	07/30/2026	Pass



Serving Size: N/A

Label Claim Profile

Analyte	Result	Specification	LOD
N-Acetyl Selank	Vial 1 = 10.14mg/vial	10mg/vial	0.01%
N-Acetyl Selank	Vial 2 = 10.17mg/vial	10mg/vial	0.01%
Identity of the analyte was confirmed by retention time concordance with a certified reference standard, per USP <621>.			
Chromatographic Purity Vial 1	>99%	>99%	0.01%
Chromatographic Purity Vial 2	>99%	>99%	0.01%
Bacterial Endotoxins (Gel-Clot: 1:10 Dilution)	<1.25 EU/ml	<5 EU/ml	$\lambda =$ 0.125 EU/mL
Rapid Sterility (Real-Time qPCR) Validated per USP <1223> in lieu of USP <71>	No Microbial DNA/RNA Detected (Pass)	No Detection	10 CFU

NOTES: Injectable peptides may include acceptable salts, sugars, or buffering agents added to improve solubility and stability. These excipients are non-chromophoric, typically not detected by HPLC, and are not considered impurities. Two vials were independently tested to evaluate batch consistency and vial-to-vial variability.

LOD = LIMIT OF DETECTION; LOQ = LIMIT OF QUANTIFICATION



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Noel Samsum
Laboratory Director
4-Aug-2026

The sample analyzed was inspected and is free from visual mold, mildew, and foreign matter. The testing procedures, equipment calibration, and maintenance are all in accordance with ISO/IEC 17025:2017 standards. The presented report is only applicable to the sample specified above and may not be applied to any similar or identical products. Reports are prohibited from being reproduced with alterations of any kind.

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Heavy Metal Analysis (USP <233>)

<u>Analyte</u>	<u>Result</u> (mcg/vial)	<u>LOQ (ppm)</u>	<u>LOD (ppm)</u>	<u>Limit</u> (mcg/day)	<u>Pass/Fail</u>
Arsenic	<LOQ	0.010	0.005	0.5	Pass
Cadmium	<LOQ	0.010	0.005	0.1	Pass
Lead	0.003	0.010	0.005	0.5	Pass
Mercury	<LOQ	0.010	0.005	0.1	Pass

Microbiological Analysis (USP <61>, USP <62>)

<u>Microbe</u>	<u>Result</u>	<u>Limit</u>	<u>Pass/Fail</u>
Total Aerobic Plate	<10 CFU/g	<100 CFU/g	Pass
Coliform	<10 CFU/g	<10 CFU/g	Pass
Bile-Tolerant Gram-Negative Bacteria	<10 CFU/g	<10 CFU/g	Pass
Yeast and Mold	<10 CFU/g	<10 CFU/g	Pass
Escherichia Coli	Negative/1g	Negative/1g	Pass
Salmonella	Negative/1g	Negative/1g	Pass
Staphylococcus Aureus	Negative/1g	Negative/1g	Pass

NOTES:

CFU = Colony Forming Unit

NS = Not Specified

NT = Not Tested

LOQ = Limit of Quantification

LOD = Limit of Detection



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