



### ORDERED BY



Your Peptide Brand

### PRODUCT

NAME VIP(10-28) 10mg  
SKU YPB.281  
CATEGORY Identity + Purity + Assay + Heavy Metals + Microbial

### SAMPLE

BATCH 20260118L04VIPS10  
TESTED 2026-04-14

DESCRIPTION Vial with white powder — sealed with plastic security cap. Flag tag attached at neck with manufacturer / product SKU number, PO, and expiration.



Submitted vial — YPB.281

AWAITING DIGITAL SIGNATURE

### ANALYTICAL RESULTS

COMPONENT	ANALYTE	SPECIFICATION	RESULT	P/F
VIP	Qualitative ID (Lambda Max)	TS λmax is a match compared to its characteristic reference standard.	Matches	✓
	Percent Purity	NLT 98% $r_{xy} = \frac{\sum (x_i - \bar{x})(y_i - \bar{y})}{\sqrt{\sum (x_i - \bar{x})^2} \cdot \sqrt{\sum (y_i - \bar{y})^2}}$	99.5 %	✓
VIP	Quantitative Assay	NLT 95% label claim $\% = \frac{A_{TS}}{A_{STD}} \times \frac{C_{STD}}{C_{TS}} \times 100$	10.41 mg	✓
	Heavy Metals (Total Quantity)	NMT 150 ppb/vial (including Pb, Cd, Hg, Ni, Fe & Co)	<10 ppb	✓
	TAMC (Aerobic/Coliform)	NMT 1,000 CFU · Incubated 48 hrs @ 37°C	0 CFU	✓
	TYMC (Yeast & Molds)	NMT 100 CFU · Incubated 48 hrs @ 30°C	0 CFU	✓



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## METHODS

- Qualitative ID - Lambda Max
- Percent Purity - Correlation Coefficient
- Quantitative Assay - Beer-Lambert
- Heavy Metals - Total Quantity
- TAMC (Total Aerobic Microbial Counts)
- TYMC (Total Yeast and Mold Counts)

Microbial specifications follow the Substances for Pharmaceutical Use compendium. UV-Vis measurements performed on a Jenway 6715 UV/Vis Spectrophotometer.



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## DISCLAIMER &amp; SCOPE OF TESTING

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NOT FOR DIAGNOSTIC, THERAPEUTIC, OR CLINICAL USE.**

The product analyzed in this Certificate of Analysis has not been evaluated by the U.S. Food and Drug Administration and is not intended to diagnose, treat, cure, or prevent any disease. Issuance of this Certificate of Analysis does not constitute FDA approval, clearance, registration, or endorsement of the product, its manufacturer, or any use thereof.

**Sample scope.**

Test results apply only to the specific sample identified by the SKU and Batch/Lot number on this report. Purity Analytics LLC tested the sample as received; chain of custody from the manufacturer to the laboratory is the manufacturer's responsibility. Subsequent handling, storage, repackaging, reconstitution, or modification by any party may alter the attributes reported here.

**Method scope.**

Pass/fail determinations reflect compliance with the acceptance criteria explicitly listed in this report and do not imply compliance with any criterion not listed. Each analytical method has inherent detection limits and measurement uncertainty; results are reported in accordance with the laboratory's validated standard operating procedures.

**No warranty.**

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