

ILS Laboratories

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Tesofensine CAPs - 500mcg

Tested for: NextGen Peptides
www.NGpeptide.com



COA #: COA-2026-YPAKKY
Lot Number: TECAP02
Accession #: ACC-2026-7142
Labeled Content: 500mcg

Analysis Date: 07/08/2026
Appearance: Good
Sample Matrix: Capsule/Tablet
Vial Size: 100 caps
Date Received: 06/30/2026

PASS



Scan to verify
authenticity at ils-lab.com
Access Code: 7GYC7J7

Method: Full QC Panel

Identity

Tesofensine CAPs

Peptide Purity

99.79%



Fentanyl
Free

Purity & Quant (HPLC)

Analyte	Specification	Result	Unit	Status
Peptide Purity (HPLC)	>= 95.0%	99.79%	%	PASS
Net Peptide Content	Report Only	0.52	mg	N/A
Identity (HPLC-RTM)	Tesofensine	Confirmed	-	PASS
Fentanyl Screen	Immunoassay, 50 ng/mL cutoff	Not Detected	-	PASS

HPLC Chromatogram



Representative chromatogram, Conformity V3

Heavy Metals (ICP-MS)

Analyte	Specification	Result	Unit	Status
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	0.433	ppm	PASS
Mercury (Hg)	<= 1.5 ppm	Not Detected	ppm	PASS
Chromium (Cr)	<= 10.0 ppm	Not Detected	ppm	PASS



Dr. Greg Kalyuzhny

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Lab Director
7/8/2026

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Verify: portal.ils-lab.com/verify/3BwWmUy7Rbtk_KD
Issued: 7/8/2026

HPLC Conformity Testing Results (4 samples tested)

Sample	Purity	NPC	ID	Result
Dedicated V0	99.79%	0.52 mg	Confirmed	PASS
Conformity V1	98.42%	0.53 mg	Confirmed	PASS
Conformity V2	98.36%	0.51 mg	Confirmed	PASS
Conformity V3	98.72%	0.51 mg	Confirmed	PASS
Mean	98.82%	0.52 mg	—	—
Std Dev	0.5750%	0.0083 mg	—	—

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	1.409 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: Purity & Quant (HPLC); Heavy Metals (ICP-MS).
- 2. The sample was confirmed to be Tesofensine CAPs 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. Chromatogram shown is representative: Conformity V3 (98.72% purity, closest to batch mean of 98.82%).




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