

ILS Laboratories

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Epithalon - 50mg



Tested for: **PeptidePharm**
PeptidePharm.com

COA #: **COA-2026-412639**
Lot Number: **ET50-05042026**
Accession #: **ACC-2026-1998**
Labeled Content: **50mg**

Method: **Full QC Panel**
Analysis Date: **05/08/2026**
Appearance: **Good**
Sample Matrix: **Lyophilized**
Date Received: **05/05/2026**

PASS



Scan to verify authenticity at ils-lab.com
Access Code: 8VUZ5KEX

Identity: **Epithalon** Peptide Purity: **99.20%**

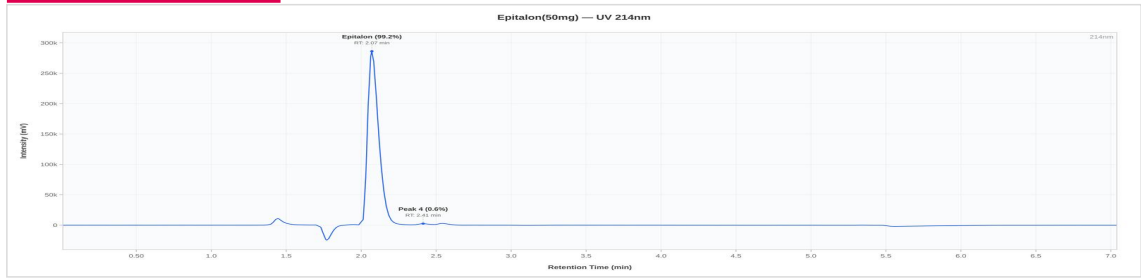
Purity & Quant (HPLC)

Analyte	Specification	Result	Unit	Status
Peptide Purity (HPLC)	>= 95.0%	99.20%	%	PASS
Net Peptide Content	Report Only	53.14	mg	N/A
Identity (HPLC-RTM)	Epithalon	Confirmed	-	PASS



Epithalon 50mg - ET50-05042026

HPLC Chromatogram



Representative chromatogram, Conformity V1 (99.06% purity, closest to batch mean of 99.13%)

HPLC Conformity Testing Results (2 samples tested)

Sample	Purity	NPC	ID	Result
Dedicated V0	99.20%	53.14 mg	Confirmed	PASS
Conformity V1	99.06%	51.47 mg	Confirmed	PASS
Mean	99.13%	52.30 mg	—	—
Std Dev	0.0700%	0.8350 mg	—	—




Dr. Greg Kalyuzhny
Lab Director
6/2/2026

COA #: **COA-2026-412639**
Access Code: **8VUZ5KEX**
Verify: portal.ils-lab.com/verify/PqR4L-GEToMxal1c
Issued: 6/2/2026

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	0.114 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: 06/02/2026. Methods: Purity & Quant (HPLC).
2. The sample was confirmed to be Epithalon 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 (99.06% purity, closest to batch mean of 99.13%).



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