

ILS Laboratories

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Melanotan I - 10

Tested for: Elevate
www.VolteraSciences.com

COA #: COA-2026-PYLY0Y
Lot Number: MT1-2026-001
Labeled Content: 10
Vial Size: 3

Analysis Date: 07/15/2026
Appearance: Good
Vial Size: 3
Date Received: 07/01/2026

Method: Full QC Panel

PASS



Scan to verify
authenticity at ils-lab.com
Access Code: EHKLS4WB

Identity	Peptide Purity	Fentanyl Free
Melanotan I	99.80%	

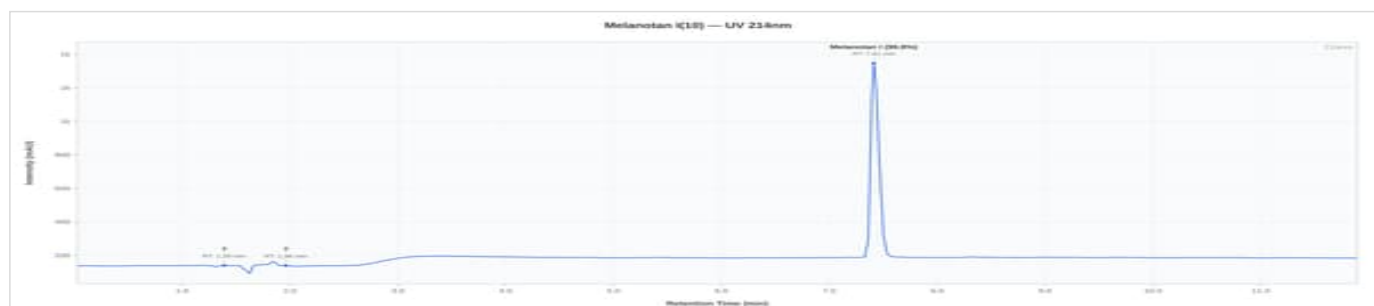
Full QC Panel

Analyte	Specification	Result	Unit	Status
Peptide Purity (HPLC)	>= 95.0%	99.80%	%	PASS
Net Peptide Content	Report Only	10.65	mg	N/A
Identity (HPLC-RTM)	Melanotan I	Confirmed	-	PASS
Identity (HPLC-RTM)	Melanotan I	Confirmed	-	PASS
Fentanyl Screen	Immunoassay, 50 ng/mL cutoff	Not Detected	-	PASS



Melanotan I 10 - MT1-2026-001

HPLC Chromatogram



Melanotan I 10 - MT1-2026-001: UV Chromatogram

Heavy Metals Analysis (ICP-MS)

Test	Specification	Result	Status
Arsenic (As)	NMT 1.5 ppm	0.897	PASS
Cadmium (Cd)	NMT 0.5 ppm	0.186	PASS
Lead (Pb)	NMT 1.0 ppm	0.193	PASS
Mercury (Hg)	NMT 1.5 ppm	0.401	PASS
Chromium (Cr)	NMT 10.0 ppm	0.388	PASS



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7/15/2026

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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	NMT 0.05 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: 07/15/2026. Methods: Full QC Panel.
2. The sample was confirmed to be Melanotan I 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.




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