

ILS Laboratories

8222 Vickers St, Suite 106, San Diego, CA 92111
(619) 329-3999 | ils-lab.com

Retatrutide - 30mg

Tested for: Voltera Sciences
www.VolteraSciences.com

COA #: COA-2026-954317
Lot Number: RT30-07082026
Accession #: ACC-2026-2048
Labeled Content: 30mg

Analysis Date: 05/08/2026
Appearance: Good
Sample Matrix: Lyophilized
Volume: 3mL
Date Received: 05/05/2026

PASS



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

Method: Full QC Panel

Identity	Peptide Purity
Retatrutide	99.62%

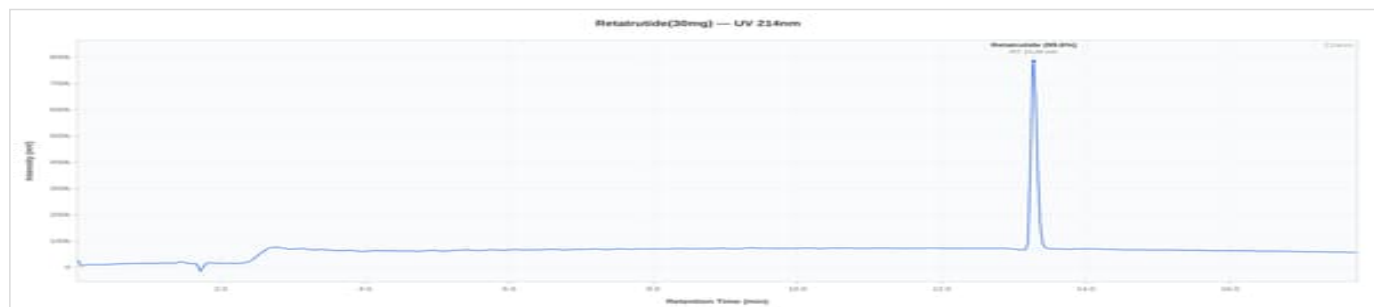
Purity, Identity & Quantitation (HPLC)

Analyte	Specification	Result	Unit	Status
Peptide Purity (HPLC)	>= 95.0%	99.62%	%	PASS
Net Peptide Content	Report Only	30.64	mg	N/A
Identity (HPLC-RTM)	Retatrutide	Confirmed	-	PASS



Retatrutide 30mg - RT30-07082026

HPLC Chromatogram



Representative chromatogram, Dedicated V0

HPLC Conformity Testing Results (2 samples tested)

Sample	Purity	NPC	ID	Result
Dedicated V0	99.62%	30.64 mg	Confirmed	PASS
Conformity V1	99.50%	31.84 mg	Confirmed	PASS
Mean	99.56%	31.24 mg	—	—
Std Dev	0.0600%	0.6000 mg	—	—

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	0.085 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

- Date Tested: 07/25/2026. Methods: Purity, Identity & Quantitation (HPLC).
- The sample was confirmed to be Retatrutide by HPLC. Identification by chromatographic retention time comparison with a reference standard.
- 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.
- Endotoxin tested per USP <85> kinetic turbidimetric method. Acceptance criteria per client specification.
- 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.
- Chromatogram shown is representative: Dedicated V0 (99.62% purity, closest to batch mean of 99.56%).

ILS Laboratories

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Retatrutide - 20mg

Tested for: Voltera Sciences
www.VolteraSciences.com

COA #: COA-2026-E3KPNQ
Lot Number: RT20-06152026
Accession #: ACC-2026-7497
Labeled Content: 20mg

Analysis Date: 07/18/2026
Appearance: Good
Sample Matrix: Lyophilized
Volume: 3mL
Date Received: 07/01/2026

PASS



Scan to verify
authenticity at ils-lab.com
Access Code: 3R5GLAKZ

Method: Full QC Panel

Identity	Peptide Purity	Fentanyl Free
Retatrutide	100.00%	

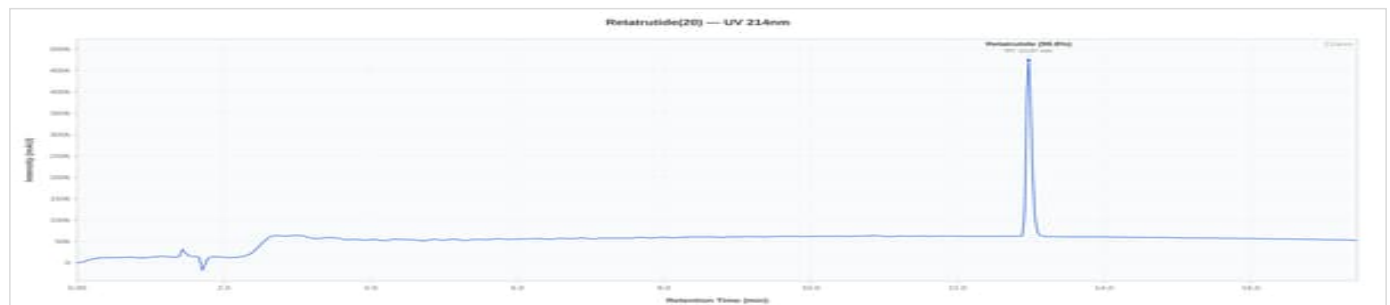
Full QC Panel

Analyte	Specification	Result	Unit	Status
Peptide Purity (HPLC)	>= 95.0%	100.00%	%	PASS
Net Peptide Content	Report Only	21.45	mg	N/A
Identity (HPLC-RTM)	Retatrutide	Confirmed	-	PASS
Fentanyl Screen	Immunoassay, 50 ng/mL cutoff	Not Detected	-	PASS



Retatrutide 20mg - RT20-06152026

HPLC Chromatogram



Representative chromatogram, Dedicated V0

HPLC Conformity Testing Results (2 samples tested)

Sample	Purity	NPC	ID	Result
Dedicated V0	99.85%	20.5 mg	Confirmed	PASS
Conformity V1	100.00%	21.45 mg	Confirmed	PASS
Mean	99.92%	20.98 mg	—	—
Std Dev	0.0750%	0.4750 mg	—	—

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	0.171 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: 07/18/2026. Methods: Full QC Panel.
2. The sample was confirmed to be Retatrutide 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: Dedicated V0 (99.85% purity, closest to batch mean of 99.92%).

ILS Laboratories

8222 Vickers St, Suite 106, San Diego, CA 92111
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VLS-3RT - 10mg

Tested for: Voltera Sciences
www.volterasciences.com

COA #: COA-2026-412743
Lot Number: RT10.05.28.26
Accession #: ACC-2026-3694
Labeled Content: 10mg

Method: Full QC Panel
Analysis Date: 06/05/2026
Appearance: Good
Vial Size: 3mL
Date Received: 06/01/2026

PASS



Scan to verify
authenticity at ils-lab.com
Access Code: NMSTTUD5

Identity	Peptide Purity	Fentanyl Free
VLS-3RT	99.15%	

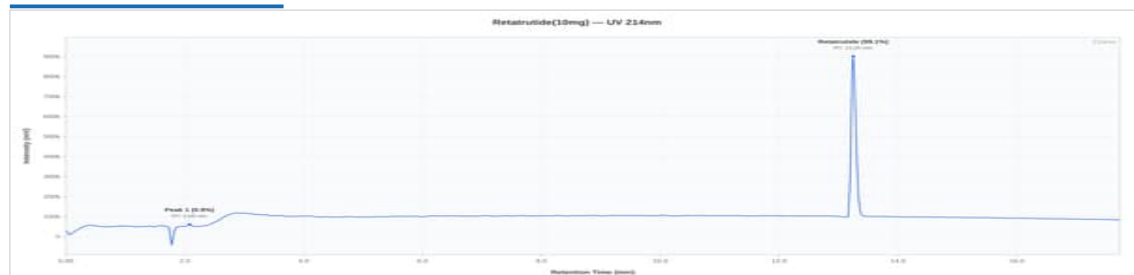
Full QC Panel

Analyte	Specification	Result	Unit	Status
Peptide Purity (HPLC)	>= 95.0%	99.15%	%	PASS
Net Peptide Content	Report Only	10.57	mg	N/A
Identity (HPLC-RTM)	Retatrutide	Confirmed	-	PASS
Fentanyl Screen	Immunoassay, 50 ng/mL cutoff	Not Detected	-	PASS



VLS-3RT 10mg - RT10.05.28.26

HPLC Chromatogram



VLS-3RT 10mg - RT10.05.28.26: UV Chromatogram

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



Dr. Greg Kalyuzhny

Dr. Greg Kalyuzhny
Lab Director
6/7/2026

COA #: COA-2026-412743
Access Code: NMSTTUD5
Verify: portal.ils-lab.com/verify/a69oQ6n129QlixIF
Issued: 6/7/2026

Endotoxin Testing (USP <85>)

Test	Specification	Result	Status
Endotoxin (USP <85>)	Report Result	0.15 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: 06/07/2026. Methods: Full QC Panel.
2. The sample was confirmed to be VLS-3RT 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.




Dr. Greg Kalyuzhny
Lab Director
6/7/2026

COA #: **COA-2026-412743**
Access Code: **NMSTTUD5**
Verify: <portal.ils-lab.com/verify/a69oQ6n129QlixIF>
Issued: 6/7/2026

ILS Laboratories

8222 Vickers St, Suite 106, San Diego, CA 92111
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VLS-3RT - 60mg



Tested for: Voltera Sciences
www.volterasciences.com

COA #: COA-2026-412744
Lot Number: RT60.05.28.26
Accession #: ACC-2026-3696
Labeled Content: 60mg

Method: Full QC Panel
Analysis Date: 06/05/2026
Appearance: Good
Vial Size: 3mL
Date Received: 06/01/2026

PASS



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authenticity at ils-lab.com
Access Code: QVRK63Y

Identity	Peptide Purity	Fentanyl Free
VLS-3RT	99.78%	

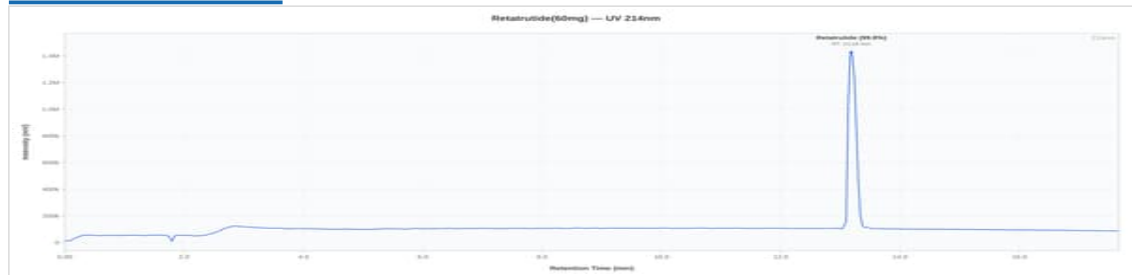
Full QC Panel

Analyte	Specification	Result	Unit	Status
Peptide Purity (HPLC)	>= 95.0%	99.78%	%	PASS
Net Peptide Content	Report Only	64.17	mg	N/A
Identity (HPLC-RTM)	Retatrutide	Confirmed	-	PASS
Fentanyl Screen	Immunoassay, 50 ng/mL cutoff	Not Detected	-	PASS



VLS-3RT 60mg - RT60.05.28.26

HPLC Chromatogram



VLS-3RT 60mg - RT60.05.28.26: UV Chromatogram

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



Dr. Greg Kalyuzhny

Dr. Greg Kalyuzhny
Lab Director
6/7/2026

COA #: COA-2026-412744
Access Code: QVRK63Y
Verify: portal.ils-lab.com/verify/nzjbtUcU08plSPYj
Issued: 6/7/2026

Endotoxin Testing (USP <85>)

Test	Specification	Result	Status
Endotoxin (USP <85>)	Report Result	0.15 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: 06/07/2026. Methods: Full QC Panel.
2. The sample was confirmed to be VLS-3RT 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.




Dr. Greg Kalyuzhny
Lab Director
6/7/2026

COA #: **COA-2026-412744**
Access Code: **QVRK63Y**
Verify: <portal.ils-lab.com/verify/nzbtUcU08pISPYj>
Issued: 6/7/2026

ILS Laboratories

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VLS-3RT - 10mg

Tested for: Voltera Sciences
www.volterasciences.com

COA #: COA-2026-412743
Lot Number: RT10.05.28.26
Accession #: ACC-2026-3694
Labeled Content: 10mg

Method: Full QC Panel
Analysis Date: 06/05/2026
Appearance: Good
Vial Size: 3mL
Date Received: 06/01/2026

PASS



Scan to verify
authenticity at ils-lab.com
Access Code: NMSTTUD5

Identity	Peptide Purity	Fentanyl Free
VLS-3RT	99.15%	

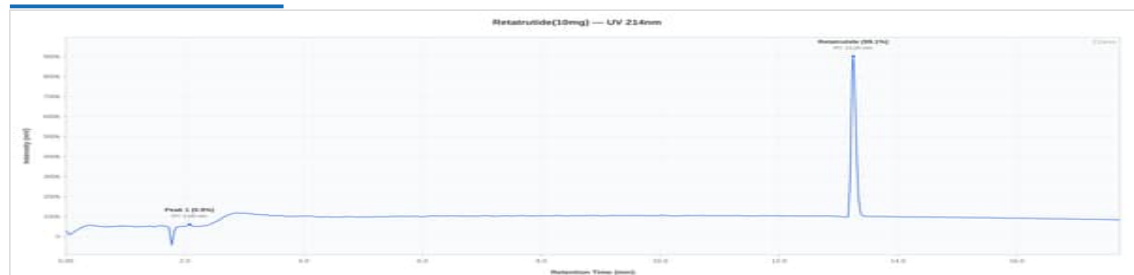
Full QC Panel

Analyte	Specification	Result	Unit	Status
Peptide Purity (HPLC)	>= 95.0%	99.15%	%	PASS
Net Peptide Content	Report Only	10.57	mg	N/A
Identity (HPLC-RTM)	Retatrutide	Confirmed	-	PASS
Fentanyl Screen	Immunoassay, 50 ng/mL cutoff	Not Detected	-	PASS



VLS-3RT 10mg - RT10.05.28.26

HPLC Chromatogram



VLS-3RT 10mg - RT10.05.28.26: UV Chromatogram

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



Dr. Greg Kalyuzhny

Dr. Greg Kalyuzhny
Lab Director
6/7/2026

COA #: COA-2026-412743
Access Code: NMSTTUD5
Verify: portal.ils-lab.com/verify/a69oQ6n129QlixIF
Issued: 6/7/2026

Endotoxin Testing (USP <85>)

Test	Specification	Result	Status
Endotoxin (USP <85>)	Report Result	0.15 EU/mL	Reported
<i>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.</i>			

Notes & Methodology

1. Date Tested: 06/07/2026. Methods: Full QC Panel.
2. The sample was confirmed to be VLS-3RT 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.




Dr. Greg Kalyuzhny
Lab Director
6/7/2026

COA #: **COA-2026-412743**
Access Code: **NMSTTUD5**
Verify: <portal.ils-lab.com/verify/a69oQ6n129QlixIF>
Issued: 6/7/2026

ILS Laboratories

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VLS-3RT - 30mg

PASS



Tested for: Voltera Sciences
www.VolteraSciences.com

COA #: COA-2026-412621
Lot Number: RT30.05.15.2026
Accession #: ACC-2026-2048
Concentration: 30mg
Sample Matrix: Lyophilized

Method: Full QC Panel
Analysis Date: 05/08/2026
Appearance: Good
Volume: 3mL
Received: 05/05/2026



Scan to verify
authenticity at ils-lab.com

Identity

VLS-3RT

Purity

99.62%

Purity & Quant (HPLC)

Analyte	Specification	Result	Unit	Status
Purity (HPLC)	>= 95.0%	99.62%	%	PASS
Net Peptide Content	Report Only	30.64	mg	N/A
Identity (ID)	Retatrutide	Confirmed	-	PASS

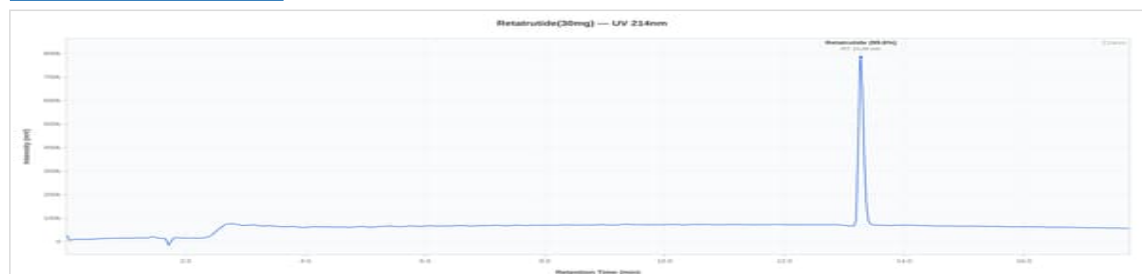


VLS-3RT 30mg - RT30.05.15.2026

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 ppm	Not Detected	ppm	PASS
Mercury (Hg)	<= 1.5 ppm	0.0887 ppm	ppm	PASS
Chromium (Cr)	<= 10 ppm	0.0693 ppm	ppm	PASS

HPLC Chromatogram



Representative chromatogram, Dedicated V0 (99.62% purity, closest to batch mean of 99.56%)



Dr. Greg Kalyuzhny

Dr. Greg Kalyuzhny
Lab Director
5/23/2026

COA #: COA-2026-412621
Access Code: VF89DCQF
Verify: portal.ils-lab.com/verify/zGQ4Breus4mhZf06
Issued: 5/23/2026



Certificate of Analysis

ILS Laboratories

8222 Vickers St, Suite 106, San Diego, CA 92111
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VLS-3RT - 30mg

Tested for: Voltera Sciences

www.VolteraSciences.com

PASS

COA #: COA-2026-412621
Lot Number: RT30.05.15.2026
Accession #: ACC-2026-2048
Concentration: 30mg
Sample Matrix: Lyophilized

Method: Full QC Panel
Analysis Date: 05/08/2026
Appearance: Good
Volume: 3mL
Received: 05/05/2026



Scan to verify
authenticity at ils-lab.com

HPLC Conformity Testing Results (2 samples tested)

Sample	Purity	NPC	ID	Result
Dedicated V0	99.62%	30.64 mg	Confirmed	PASS
Conformity V1	99.50%	31.84 mg	Confirmed	PASS
Mean	99.56%	31.24 mg	—	—
Std Dev	0.0600%	0.6000 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	0.085 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

- Date Tested: 05/23/2026. Methods: Purity & Quant (HPLC); Heavy Metals (ICP-MS).
- The sample was confirmed to be VLS-3RT by HPLC. Identification by chromatographic retention time comparison with a reference standard.
- Endotoxin acceptance limits are product-specific. Results reported for client evaluation.
- Chromatogram shown is representative: Dedicated V0 (99.62% purity, closest to batch mean of 99.56%).



Dr. Greg Kalyuzhny

Dr. Greg Kalyuzhny
Lab Director
5/23/2026

COA #: COA-2026-412621
Access Code: VF89DCQF
Verify: portal.ils-lab.com/verify/zGQ4Breus4mhzF06
Issued: 5/23/2026

Client: Voltera

Certified: 05/14/2026

This Certificate of Analysis certifies that the sample listed herein was tested by Kovera Labs using validated analytical methods and was found to meet the stated specifications at the time of analysis.

SAMPLE INFORMATION

Product	VLS-3RT	Form	Lyophilized powder
Batch	VLSRT.05.02.2026	Labeled Qty	10 mg
Mol. Formula	C221H342N46O68	CAS Number	2381089-83-2
Cap Color	Yellow	Crimp Color	Silver

TEST RESULTS

	REFERENCE STANDARD	RESULT
Batch Avg Purity	(>98%)	99.951%
Batch Avg Net Content	(10mg ± 10%)	11.36 mg
Identity Confirmation (LC-MS)	(VLS-3RT)	VLS-3RT
Endotoxin Safety Screen	(≤0.5 EU/mL)	PASS
Microbial Sterility Screen	(No Growth)	No Growth

HEAVY METAL SCREENING

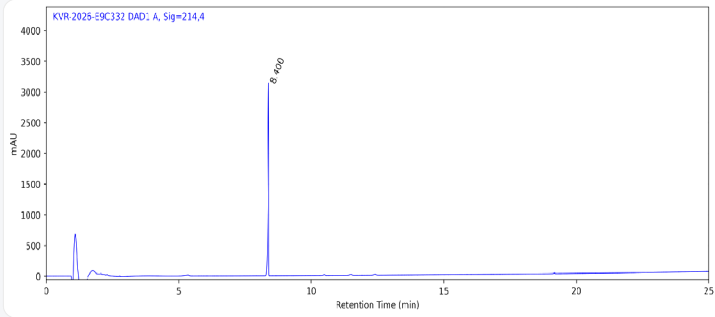
Analyte	Result	Status
Arsenic (As)	Negative	
Cadmium (Cd)	Negative	
Lead (Pb)	Negative	
Mercury (Hg)	Negative	

BATCH CONFORMITY RESULTS

Vial	Purity (%)	Net Content (mg)
Vial 1	99.939	11.39
Vial 2	99.962	11.32
Batch Average	99.951%	11.36 mg

CHROMATOGRAM

Method: RP-HPLC | Column: C18 | Detection: DAD @ 214 nm



Lemar Arghandival
Lab Director



koveralabs.com/verify

Report#: KVR-2026-0F26E3
Access Code: XZ3WOJX

Client: Voltera

Certified: 05/14/2026

This Certificate of Analysis certifies that the sample listed herein was tested by Kovera Labs using validated analytical methods and was found to meet the stated specifications at the time of analysis.

SAMPLE INFORMATION

Product	VLS-3RT	Form	Lyophilized powder
Batch	VLSRT.05.02.2026	Labeled Qty	20 mg
Mol. Formula	C225H348N48O68	CAS Number	2023788-19-2
Cap Color	White	Crimp Color	Silver

TEST RESULTS

	REFERENCE STANDARD	RESULT
Batch Avg Purity	(>98%)	99.864%
Batch Avg Net Content	(20mg ± 10%)	21.31 mg
Identity Confirmation (LC-MS)	(VLS-3RT)	VLS-3RT
Endotoxin Safety Screen	(≤0.5 EU/mL)	PASS
Microbial Sterility Screen	(No Growth)	No Growth

HEAVY METAL SCREENING

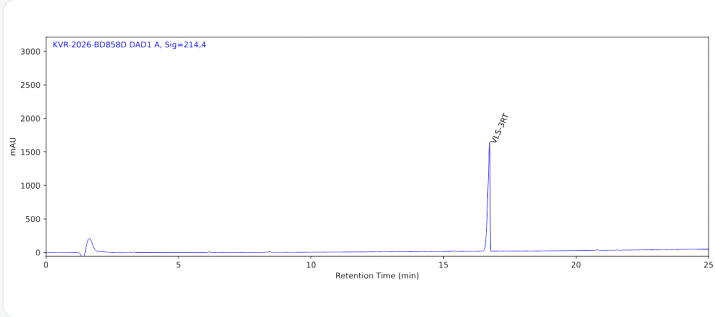
Analyte	Result	Status
Arsenic (As)	Negative	
Cadmium (Cd)	Negative	
Lead (Pb)	Negative	
Mercury (Hg)	Negative	

BATCH CONFORMITY RESULTS

Vial	Purity (%)	Net Content (mg)
Vial 1	99.869	21.31
Vial 2	99.858	21.31
Batch Average	99.864%	21.31 mg

CHROMATOGRAM

Method: RP-HPLC | Column: C18 | Detection: DAD @ 214 nm



Lemar Arghandival
Lab Director



koveralabs.com/verify

Report#: KVR-2026-BD858D
Access Code: IK6M3RH

Client: Voltera

Certified: 05/17/2026

This Certificate of Analysis certifies that the sample listed herein was tested by Kovera Labs using validated analytical methods and was found to meet the stated specifications at the time of analysis.

SAMPLE INFORMATION

Product	VLS-3RT	Form	Lyophilized powder
Batch	VLSRT.05.17.2026	Labeled Qty	20 mg
Mol. Formula	C221H342N46O68	CAS Number	2381089-83-2
Cap Color	Black	Crimp Color	Silver

TEST RESULTS

	REFERENCE STANDARD	RESULT
Batch Avg Purity	(>98%)	99.862%
Batch Avg Net Content	(20mg ± 10%)	22.88 mg
Identity Confirmation (LC-MS)	(VLS-3RT)	VLS-3RT
Endotoxin Safety Screen	(≤0.5 EU/mL)	PASS
Microbial Sterility Screen	(No Growth)	No Growth

HEAVY METAL SCREENING

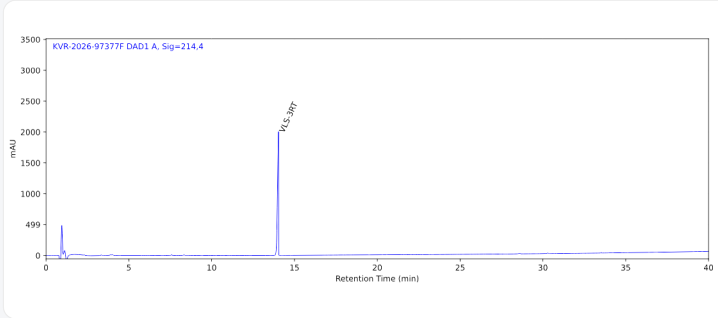
Analyte	Result	Status
Arsenic (As)	Negative	
Cadmium (Cd)	Negative	
Lead (Pb)	Negative	
Mercury (Hg)	Negative	

BATCH CONFORMITY RESULTS

Vial	Purity (%)	Net Content (mg)
Vial 1	99.821	22.87
Vial 2	99.907	22.83
Vial 3	99.858	22.94
Batch Average	99.862%	22.88 mg

CHROMATOGRAM

Method: RP-HPLC | Column: C18 | Detection: DAD @ 214 nm



Amir Arghandinal
Lab Director



koveralabs.com/verify

Report#: KVR-2026-97377F
Access Code: UH9W2WI