



CERTIFICATE OF ANALYSIS

Bacteriostatic Water

3 ml

LOT NUMBER
APX-2026-0511-B

REPORT NUMBER
APX-COA-2026-0511-B

COA ISSUE DATE
2026-05-11

COMPOUND REFERENCE

Material family
solvent or reagent
Material form
aqueous liquid in vial

MATERIAL & COMPOSITION

Nominal component amounts
Benzyl alcohol: 27 mg
Recorded storage
2-8 C, protected from light

Reagent quality measurements

pH of native liquid
5.82 pH

4.61 |-----| 6.99

Criterion: 5.1 to 6.5

Preservative efficacy bacteria day 14 reduction
3.6 log₁₀ reduction

2.79 |-----| 3.81

Criterion: ≥ 3

Criterion uses the result unit. Independent expanded scales.

Measured concentration

Benzyl alcohol concentration
9.00414 mg/mL

8.694 |-----| 9.306

Criterion: 8.82 to 9.18

Criterion uses the result unit. Independent expanded scales.

59 recorded endpoints across 21 test groups. Results apply to this configuration and lot. For research use only; not for human or animal use.

APPROVED BY

K. Norwood

COA issued 2026-05-11

ONLINE REPORT

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02 Contamination & microbiology
Metals, solvents and microbial results

03+ Complete assay results
Methods, specifications and analytical context

Contamination & microbiology

All applicable tested criteria met

32 applicable endpoints reviewed against their stated specifications.

ELEMENTAL IMPURITIES

Lead	✓	PASS
Cadmium	✓	PASS
Arsenic	✓	PASS
Mercury	✓	PASS
Cobalt	✓	PASS
Vanadium	✓	PASS
Nickel	✓	PASS
Palladium	✓	PASS
Platinum	✓	PASS
Rhodium	✓	PASS
Ruthenium	✓	PASS
Iridium	✓	PASS

RESIDUAL SOLVENTS

Methanol	✓	PASS
Acetonitrile	✓	PASS
Dichloromethane	✓	PASS
N, N-dimethylformamide	✓	PASS
Ethanol	✓	PASS
Acetone	✓	PASS
Ethyl acetate	✓	PASS
Toluene	✓	PASS
n-Hexane	✓	PASS
Benzene	✓	PASS

MICROBIOLOGY & ENDOTOXINS

Sterility: FTM	✓	PASS
Sterility: SCDM	✓	PASS
Bacterial endotoxins	✓	PASS
TAMC	✓	PASS
TYMC	✓	PASS
Escherichia coli	✓	PASS
Staphylococcus aureus	✓	PASS
Pseudomonas aeruginosa	✓	PASS
Salmonella species	✓	PASS
Candida albicans	✓	PASS

PASS indicates that the recorded result meets its stated criterion for the tested sample. Exact values, detection limits, sampling scope and methods follow in the complete results. For research use only.

Complete assay results

Reported values, units and acceptance outcomes are reproduced separately for each endpoint. Method keys link to the method reference section.

Endpoint	Reported result / basis	Acceptance criterion	Method / outcome
T01 / Identity and structure			
01 / Orthogonal identity against specified reference	Conforms to the assigned reference; see identity basis Unit: qualitative Basis B01	Conforms to the specified acceptance criterion Unit: qualitative	M01 PASS C01
02 / Orthogonal fingerprint similarity	= 0.9994 similarity (0-1) Basis B02	>= 0.995 Unit: similarity (0-1)	M01 PASS C02
T02 / Chromatographic purity			
03 / Applicability decision	Not applicable — Bulk water or dilute aqueous reagent is controlled by calibrated concentration and reagent tests; 99.7% finished-solution HPLC area purity is not meaningful. Unit: qualitative Basis B03	Same as reported result Unit: qualitative Basis B04	M02 N/A — justified C03
T03 / Active content and assay			
05 / Benzyl alcohol active content	= 27.0124 mg/vial Numeric value: 27.01242 Basis B06	26.46 to 27.54 Unit: mg/vial	M04 PASS C02
06 / Benzyl alcohol label claim assay	= 100.046 % label claim Basis B07	98 to 102 Unit: % label claim	M04 PASS C02
07 / Benzyl alcohol concentration	= 9.00414 mg/mL Basis B08	8.82 to 9.18 Unit: mg/mL	M04 PASS C02
T04 / Blend composition			
10 / Applicability decision	Not applicable — One declared active or a water/preservative reagent. Preservative is separately quantified under T14; there is no multi-active blend ratio. Unit: qualitative Basis B03	Same as reported result Unit: qualitative Basis B04	M06 N/A — justified C03
T05 / Related substances and degradants			
04 / Related organic impurity response in reagent assay	= 0.025 % of acetic acid or benzyl alcohol analyte response Basis B05	<= 0.1 Unit: % of acetic acid or benzyl alcohol analyte response	M03 PASS C04

Endpoint	Reported result / basis	Acceptance criterion	Method / outcome
T06 / Water content			
12 / Applicability decision	Not applicable — Water is the formulation vehicle; residual-moisture percentage would misdescribe an aqueous formulation. Vehicle quantity is specified separately. Unit: qualitative Basis B03	Same as reported result Unit: qualitative Basis B04	M07 N/A — justified C03
T07 / Counterions and associated species			
13 / Applicability decision	Not applicable — No fixed counterion salt is assigned in this formulation; vehicle ions and intentional metal complexes are treated by their component specifications. Unit: qualitative Basis B03	Same as reported result Unit: qualitative Basis B04	M08 N/A — justified C03
T08 / Residual solvents			
14 / Methanol	= 19.428 ug/mL formulation Basis B11	<= 3000 Unit: ug/mL formulation	M09 PASS C06
15 / Acetonitrile	= 7.722 ug/mL formulation Basis B11	<= 410 Unit: ug/mL formulation	M09 PASS C07
16 / Dichloromethane	= 3.701 ug/mL formulation Basis B11	<= 600 Unit: ug/mL formulation	M09 PASS C08
17 / N, N-dimethylformamide	= 4.444 ug/mL formulation Basis B11	<= 880 Unit: ug/mL formulation	M09 PASS C09
18 / Ethanol	= 38.412 ug/mL formulation Basis B11	<= 5000 Unit: ug/mL formulation	M09 PASS C06
19 / Acetone	= 11.919 ug/mL formulation Basis B11	<= 5000 Unit: ug/mL formulation	M09 PASS C09
20 / Ethyl acetate	= 9.358 ug/mL formulation Basis B11	<= 5000 Unit: ug/mL formulation	M09 PASS C09
21 / Toluene	= 1.856 ug/mL formulation Basis B11	<= 890 Unit: ug/mL formulation	M09 PASS C10
22 / n-Hexane	= 1.173 ug/mL formulation Basis B11	<= 290 Unit: ug/mL formulation	M09 PASS C10

Endpoint	Reported result / basis	Acceptance criterion	Method / outcome
23 / Benzene	= 0.072 ug/mL formulation Basis B11	<= 2 Unit: ug/mL formulation	M09 PASS C11
T09 / Elemental impurities			
24 / Lead	= 0.0126 ug/mL formulation Basis B12	<= 0.5 Unit: ug/mL formulation	M10 PASS C12
25 / Cadmium	= 0.0081 ug/mL formulation Basis B12	<= 0.2 Unit: ug/mL formulation	M10 PASS C12
26 / Arsenic	= 0.0182 ug/mL formulation Basis B12	<= 0.5 Unit: ug/mL formulation	M10 PASS C12
27 / Mercury	= 0.0061 ug/mL formulation Basis B12	<= 0.1 Unit: ug/mL formulation	M10 PASS C12
28 / Cobalt	= 0.0214 ug/mL formulation Basis B12	<= 1 Unit: ug/mL formulation	M10 PASS C12
29 / Vanadium	= 0.0225 ug/mL formulation Basis B12	<= 1 Unit: ug/mL formulation	M10 PASS C12
30 / Nickel	= 0.0382 ug/mL formulation Basis B12	<= 2 Unit: ug/mL formulation	M10 PASS C12
31 / Palladium	= 0.012 ug/mL formulation Basis B12	<= 1 Unit: ug/mL formulation	M10 PASS C12
32 / Platinum	= 0.0093 ug/mL formulation Basis B12	<= 1 Unit: ug/mL formulation	M10 PASS C12
33 / Rhodium	= 0.0074 ug/mL formulation Basis B12	<= 1 Unit: ug/mL formulation	M10 PASS C12
34 / Ruthenium	= 0.009 ug/mL formulation Basis B12	<= 1 Unit: ug/mL formulation	M10 PASS C12
35 / Iridium	= 0.0055 ug/mL formulation Basis B12	<= 1 Unit: ug/mL formulation	M10 PASS C12

Endpoint	Reported result / basis	Acceptance criterion	Method / outcome
T10 / Sterility test			
42 / Sterility: FTM	No growth at 14 days in all 20 units Unit: qualitative Basis B19	Conforms to the specified acceptance criterion Unit: qualitative	M14 PASS C13
43 / Sterility: SCDM	No growth at 14 days in all 20 units Unit: qualitative Basis B20	Conforms to the specified acceptance criterion Unit: qualitative	M14 PASS C13
T11 / Bacterial endotoxins			
44 / Bacterial endotoxins	<0.025 EU/mL Basis B21	<= 0.1 Unit: EU/mL	M15 PASS C14
T12 / Microbial enumeration			
45 / TAMC	<1 CFU/mL Basis B22	<= 100 Unit: CFU/mL	M16 PASS C15
46 / TYMC	<1 CFU/mL Basis B22	<= 10 Unit: CFU/mL	M16 PASS C15
47 / Escherichia coli	Not detected in 1 mL Unit: qualitative Basis B23	Conforms to the specified acceptance criterion Unit: qualitative	M16 PASS C16
48 / Staphylococcus aureus	Not detected in 1 mL Unit: qualitative Basis B23	Conforms to the specified acceptance criterion Unit: qualitative	M16 PASS C16
49 / Pseudomonas aeruginosa	Not detected in 1 mL Unit: qualitative Basis B23	Conforms to the specified acceptance criterion Unit: qualitative	M16 PASS C16
50 / Salmonella species	Not detected in 1 mL Unit: qualitative Basis B23	Conforms to the specified acceptance criterion Unit: qualitative	M16 PASS C16
51 / Candida albicans	Not detected in 1 mL Unit: qualitative Basis B23	Conforms to the specified acceptance criterion Unit: qualitative	M16 PASS C16
T13 / pH and analytical solution			
36 / pH of native liquid	= 5.82 pH Basis B13	5.1 to 6.5 Unit: pH	M11 PASS C02
T14 / Excipients and preservatives			
37 / Benzyl alcohol preservative	= 9.00414 mg/mL Basis B14	8.55 to 9.45 Unit: mg/mL	M12 PASS C02

Endpoint	Reported result / basis	Acceptance criterion	Method / outcome
38 / Preservative efficacy bacteria day 14 reduction	= 3.6 log10 reduction Basis B15	>= 3 Unit: log10 reduction	M12 PASS C02
39 / Preservative efficacy fungi day 28 growth	No increase from inoculum Unit: qualitative Basis B16	Conforms to the specified acceptance criterion Unit: qualitative	M12 PASS C01
T15 / Unit uniformity			
08 / Benzyl alcohol individual-unit mean	= 100.046 % label claim Basis B03	98 to 102 Unit: % label claim Basis B04	M05 PASS C05
09 / Benzyl alcohol individual-unit RSD	= 0.3523 % Basis B09	<= 2 Unit: %	M05 PASS C05
11 / Mean extractable volume	= 3 mL/container Basis B10	2.97 to 3.06 Unit: mL/container	M05 PASS C05
T16 / Biological activity			
52 / Applicability decision	Not applicable — No qualified, reference-traceable potency endpoint is established for this selected identity. Identity/content data cannot justify an invented biological potency value; exploratory biological activity would require a separately defined assay. Unit: qualitative Basis B03	Same as reported result Unit: qualitative Basis B04	M17 N/A — justified C03
T17 / Protein and conjugate characterization			
53 / Applicability decision	Not applicable — No recombinant protein, polymer conjugate or complex mixture for this material. Defined synthetic peptide identity is characterized under T01; GHK copper speciation, when relevant, is explicit. Unit: qualitative Basis B03	Same as reported result Unit: qualitative Basis B04	M18 N/A — justified C03
T18 / Process impurities			
54 / Applicability decision	Not applicable — chemical synthesis without cell/tissue manufacturing inputs; HCP, host DNA and Protein A are not relevant. Chemical process impurities are covered under T05/T08/T09. Unit: qualitative Basis B03	Same as reported result Unit: qualitative Basis B04	M19 N/A — justified C03

Endpoint	Reported result / basis	Acceptance criterion	Method / outcome
T19 / Particles and container closure			
55 / Subvisible particles $\geq 10\mu\text{m}$	= 21 particles/container Basis B24	≤ 6000 Unit: particles/container	M20 PASS C17
56 / Subvisible particles $\geq 25\mu\text{m}$	= 2 particles/container Basis B24	≤ 600 Unit: particles/container	M20 PASS C17
57 / Vacuum decay package integrity	10/10test closures conform; positive-leak controls detected Unit: qualitative Basis B25	Conforms to the specified acceptance criterion Unit: qualitative	M20 PASS C18
T20 / Stability observations			
58 / Reported 6 month assay retention	= 99.91 % of initial assay Basis B26	98 to 102 Unit: % of initial assay	M21 PASS C02
59 / Dating disposition	6 month review point; actual expiry/retest not assigned Unit: qualitative Basis B26	Conforms to the specified acceptance criterion Unit: qualitative	M21 PASS C01
T21 / Appearance and physical inspection			
40 / Appearance	colorless; clear solution Unit: qualitative Basis B17	Conforms to the specified acceptance criterion Unit: qualitative	M13 PASS C01
41 / Visible foreign matter	None observed in 3 inspected units Unit: qualitative Basis B18	Conforms to the specified acceptance criterion Unit: qualitative	M13 PASS C01

Method reference

The keys below identify the methods referenced in the result tables.

Key / method ID	Method	Technique
M01 M-T01-DEFAULT	Orthogonal chemical/structural identity	LC-HRMS/MS plus sequence/connectivity-sensitive reference comparison
M02 M-T02-DEFAULT	Chromatographic purity	RP-UPLC DAD; SEC for protein size species; HILIC/CAD for nonchromophoric analytes
M03 M-T05-DEFAULT	Related substances	Stability-indicating LC integration; orthogonal size/charge separation for biologics
M04 M-T03-DEFAULT	Calibrated material content	Independent quantitative LC/CAD, protein assay or bioactivity calibration for IU
M05 M-T15-DEFAULT	Individual-unit uniformity	Ten separate unit preparations and fill measurements
M06 M-T04-DEFAULT	Component-resolved blend assay	Separate calibrated selective assay for every declared component
M07 M-T06-DEFAULT	Water	Coulometric Karl Fischer
M08 M-T07-DEFAULT	Counterion composition	Ion chromatography and stoichiometric accounting
M09 M-T08-DEFAULT	Residual solvent panel	headspace GC-MS
M10 M-T09-DEFAULT	Elemental impurity panel	ICP-MS
M11 M-T13-DEFAULT	pH and solution state	Calibrated potentiometry
M12 M-T14-DEFAULT	Excipients/preservatives	Component-specific quantitative LC/CAD or formulation identity
M13 M-T21-DEFAULT	Appearance	Controlled visual examination
M14 M-T10-DEFAULT	Sterility observations in two media with matrix-specific preparation	Membrane filtration/rinsing or neutralized direct inoculation for Lemon dispersion
M15 M-T11-DEFAULT	Bacterial endotoxins	Recombinant factor C fluorometric assay
M16 M-T12-DEFAULT	Microbial enumeration/organisms	Membrane filtration enumeration and enrichment
M17 M-T16-DEFAULT	Functional biological activity	Reference-relative concentration-response assay
M18 M-T17-DEFAULT	Protein/conjugate characterization	SEC-MALS, CE-SDS, peptide mapping, glycan/PEG profiling

Key / method ID	Method	Technique
M19 M-T18-DEFAULT	Biological process impurity panel	Host/process-specific ELISA and qPCR
M20 M-T19-DEFAULT	Particles and package integrity	Light obscuration plus vacuum decay
M21 M-T20-DEFAULT	Stability observation	Time-zero/1/3/6-month trend

Analytical detail appendix

Basis keys preserve the exact analytical scope of each result. Detection limits and sampling context are reported separately from measured values. N/A limits are not numerical detection limits and are omitted from the context key.

Result and criterion bases

Key	Basis
B01	bacteriostatic laboratory water:benzyl alcohol9.00 mg/mL(0.900%w/v) in qualified water; 3.00 mL and 10.00 mL nominal fills. No sodium chloride. Sterility/endotoxin scope is separately declared.
B02	spectrum/peptide-map comparison; score is not proof of stereochemistry
B03	specified sample/form only
B04	material-specific limit
B05	Analyte-specific reagent assay; not water-purity percentage
B06	Assigned component material basis; excludes vehicle/excipient/water; salt basis explicitly in identity specification
B07	Independent calibrated amount / nominal amount x 100; not chromatographic area purity
B08	Amount per nominal 3 mL fill
B09	Precision criterion for the listed specification, independently of AV
B10	fill specification; no injection volume recommendation
B11	Material-based limit. Liquid numerical limits are listed specification and not ICH Option1 mass ppm.
B12	Total element; no speciation. material-based criterion, not a dose/route-derived safety limit.
B13	As supplied,25.0C; buffer-solution compatibility is not the pH of dry material
B14	0.90%w/v target=9.00 mg/mL; same independently calibrated data as T03
B15	Additional challenge study; separate from preservative concentration
B16	Separate C.albicans and A.brasiliensis observations in challenge data
B17	Three containers under diffuse white illumination
B18	Visual observation only; subvisible particles reported separately
B19	FTM30-35C; tested units only; growth-promotion/suitability controls conform
B20	SCDM20-25C; tested units only; no sterility assurance level assigned
B21	Sample reporting limit after1:5 dilution; threshold is and not based on administration route/dose
B22	Zero colonies observed from1g or1 mL equivalent filter; report <1, not exact absence
B23	Specified-organism enrichment in explicit1g/1 mL scope; not a universal pharmacopeial panel
B24	As-supplied liquid or entire dry vial dissolved to stated analytical volume; limits, no injectable compliance claim
B25	method vacuum-decay threshold0.20kPa/60s;5um positive defect control. No universal leakage-to-sterility equivalence.
B26	Reported change relative to the initial assay. This report does not assign a retest date or shelf life.

Sampling and detection context

Key	Reported context
C01	Analytical replicates: 2; Statistic type: individual/qualitative observation; Units sampled: 3; Units type: finished containers
C02	Analytical replicates: 2; Statistic type: mean of independent sample preparations; Units sampled: 3; Units type: finished containers
C03	Analytical replicates: 0; Statistic type: individual/qualitative observation; Units sampled: 0; Units type: finished containers
C04	Analytical replicates: 2; Statistic type: mean of independent sample preparations; Units sampled: 3; Units type: finished containers; LOD: 0.002 % of acetic acid or benzyl alcohol analyte response; LOQ: 0.005 % of acetic acid or benzyl alcohol analyte response
C05	Analytical replicates: 1; Statistic type: mean of independent sample preparations; Units sampled: 10; Units type: finished containers
C06	Analytical replicates: 2; Statistic type: mean of independent sample preparations; Units sampled: 3; Units type: finished containers; LOD: 1 ug/mL formulation; LOQ: 3 ug/mL formulation
C07	Analytical replicates: 2; Statistic type: mean of independent sample preparations; Units sampled: 3; Units type: finished containers; LOD: 0.3 ug/mL formulation; LOQ: 1 ug/mL formulation
C08	Analytical replicates: 2; Statistic type: mean of independent sample preparations; Units sampled: 3; Units type: finished containers; LOD: 0.2 ug/mL formulation; LOQ: 0.6 ug/mL formulation
C09	Analytical replicates: 2; Statistic type: mean of independent sample preparations; Units sampled: 3; Units type: finished containers; LOD: 0.5 ug/mL formulation; LOQ: 1.5 ug/mL formulation
C10	Analytical replicates: 2; Statistic type: mean of independent sample preparations; Units sampled: 3; Units type: finished containers; LOD: 0.1 ug/mL formulation; LOQ: 0.3 ug/mL formulation
C11	Analytical replicates: 2; Statistic type: mean of independent sample preparations; Units sampled: 3; Units type: finished containers; LOD: 0.01 ug/mL formulation; LOQ: 0.03 ug/mL formulation
C12	Analytical replicates: 2; Statistic type: mean of independent sample preparations; Units sampled: 3; Units type: finished containers; LOD: 0.0005 ug/mL formulation; LOQ: 0.002 ug/mL formulation
C13	Analytical replicates: 1; Statistic type: individual/qualitative observation; Units sampled: 20; Units type: finished containers
C14	Analytical replicates: 2; Statistic type: mean of independent sample preparations; Units sampled: 3; Units type: finished containers; LOD: 0.01 EU/mL; LOQ: 0.025 EU/mL
C15	Analytical replicates: 2; Statistic type: mean of independent sample preparations; Units sampled: 3; Units type: finished containers; LOD: 1 CFU/mL; LOQ: 1 CFU/mL
C16	Analytical replicates: 1; Statistic type: individual/qualitative observation; Units sampled: 1; Units type: finished containers
C17	Analytical replicates: 1; Statistic type: mean of independent sample preparations; Units sampled: 3; Units type: finished containers
C18	Analytical replicates: 1; Statistic type: individual/qualitative observation; Units sampled: 10; Units type: finished containers

Supporting analytical details

01 / Orthogonal identity against specified reference

Field	Reported value	Unit / note
identified structure or sequence	bacteriostatic laboratory water:benzyl alcohol9.00 mg/mL(0.900%w/v) in qualified water; 3.00 mL and 10.00 mL nominal fills. No sodium chloride. Sterility/endotoxin scope is separately declared.	text
neutral average mass	Not assigned: construct/distribution or no supported unique species	Da or explicit reason
neutral monoisotopic mass	Not assigned — see component-specific HRMS records; a reference mass is not a measured neutral mass	Explicit context; see value Method and sample context; reference facts are in Identity Reference.
observed mz	Not assigned — heterogeneous construct or no defined ion; orthogonal identity documented instead	Explicit context; see value Method and sample context; reference facts are in Identity Reference.
mass tolerance	+/-5 ppm for an explicitly defined monoisotopic ion; not 5 Da.	Explicit context; see value Method and sample context; reference facts are in Identity Reference.
reduced deglycosylated or intact	Intact specified parent for defined compounds; glycoproteins and PEG retain the specified population heterogeneity. No single exact intact glycoprotein mass is invented.	Explicit context; see value Method and sample context; reference facts are in Identity Reference.
stereochemistry method	Component-specific chiral amino acid analysis or orthogonal structure comparison, where applicable; HRMS cannot distinguish stereoisomers.	Explicit context; see value Method and sample context; reference facts are in Identity Reference.
identity standard traceability	RS-2567; internal standard, not a purchased certified reference material	text

05 / Benzyl alcohol active content

Field	Reported value	Unit / note
sample preparation volume	27	mL
dilution factor	10	fold

08 / Benzyl alcohol individual-unit mean

Field	Reported value	Unit / note
physical unit count	1	Explicit context; see value Method and sample context; reference facts are in Identity Reference.

44 / Bacterial endotoxins

Field	Reported value	Unit / note
dilution factor	5	Explicit context; see value Method and sample context; reference facts are in Identity Reference.
maximum valid dilution if used	20	Explicit context; see value Method and sample context; reference facts are in Identity Reference.

55 / Subvisible particles >= 10um

Field	Reported value	Unit / note
packaging sample count	10	Explicit context; see value Method and sample context; reference facts are in Identity Reference.

END OF REPORT

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