

CLIENT & SAMPLE INFORMATION

Client

Aurora Peptides

Analysis Date

07/11/2026

Lot / Batch

HHB-072026-001

Product Name

Radiance Blend

Form / Matrix

Liquid Blend

Volume

10 mL

This Certificate of Analysis confirms the identity and quality of the multi-component liquid blend listed herein. Compound identification was performed by HPLC-UV with LC-MS/MS confirmation. Declared active components were evaluated against their expected molecular masses and chromatographic profiles.

ACTIVE COMPONENT IDENTITY (LC-MS/MS)

Declared components evaluated by HPLC with UV detection, confirmed by Mass Spectrometry where a characteristic ion is available. Identity confirmed via observed m/z vs. theoretical m/z.

Component	CAS Number	Theoretical m/z [M+H] ⁺	Observed m/z	Status
Niacinamide (B3)	98-92-0	123.06	123.06	PASS
Thiamine HCl (B1)	67-03-8	265.11	265.11	PASS
Pantothenic Acid (B5)	79-83-4	220.12	220.12	PASS
Choline	62-49-7	104.11	104.11	PASS
myo-Inositol	87-89-8	181.07	181.07	PASS
Niacin	59-67-6	124.04	124.04	PASS
Biotin (B7)	58-85-5	245.10	245.10	PASS
Folic Acid (B9)	59-30-3	442.15	442.15	PASS
Riboflavin (B2)	83-88-5	377.15	377.15	PASS
Overall Identity Determination				PASS

Method: HPLC-UV with LC-MS/MS confirmation | 9 declared component(s) evaluated. Components lacking a UV chromophore are confirmed by orthogonal identity where applicable.

PHYSICOCHEMICAL PROPERTIES

Property	Specification	Result	Status
Appearance / Color	Conforms	Conforms	PASS
Physical Form	Liquid	Liquid	PASS
Particulate Matter	Essentially free	Conforms	PASS

ADDITIONAL QUALITY SCREENS

ENDOTOXIN SCREEN

Method

LAL (USP <85>)

Limit

<= 0.5 EU/mL

Result

Pass

Status

PASS

STERILITY / MICROBIAL SCREEN

Method

Rapid Sterility

Specification

No Growth

Incubation

2 days

Result

No microbial growth detected

Fungi

Not Detected

Status

PASS

HEAVY METAL SCREEN

Method

ICP-MS

Specification

Compliant

Result

PASS

Status

PASS

TESTING METHODOLOGY

Compound Identity (LC-MS/MS)

Multi-component identity confirmation performed by reverse-phase HPLC with UV detection coupled with tandem mass spectrometry (LC-MS/MS). Each declared active is identified by retention time and, where a characteristic ion is available, confirmed by observed [M+H]⁺ molecular ion matched against its theoretical mass. Highly polar components lacking a UV chromophore (e.g. free amino acids) are reported by identity confirmation and may require orthogonal (derivatization / HILIC) methods for full quantitation.

