

CLIENT & SAMPLE INFORMATION

|               |                  |                |              |
|---------------|------------------|----------------|--------------|
| Client        | Evolabs Research | Certified Date | 07/22/2026   |
| Lot / Batch   | 7312026          | Product Name   | Lipo-C + B12 |
| 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                      | Theoretical m/z [M+H] <sup>+</sup> | Observed m/z | Status |
|--------------------------------|------------------------------------|--------------|--------|
| L-Methionine                   | 150.06                             | 150.06       | PASS   |
| Inositol                       | 181.07                             | 181.07       | PASS   |
| Choline                        | 104.11                             | 104.11       | PASS   |
| L-Carnitine                    | 162.11                             | 162.11       | PASS   |
| Vitamin B12                    | —                                  |              | PASS   |
| Overall Identity Determination |                                    |              | PASS   |

Method: HPLC-UV with LC-MS/MS confirmation | 5 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 | < 0.10 EU/mL   |
| 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]<sup>+</sup> 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.

