

CERTIFICATE OF ANALYSIS

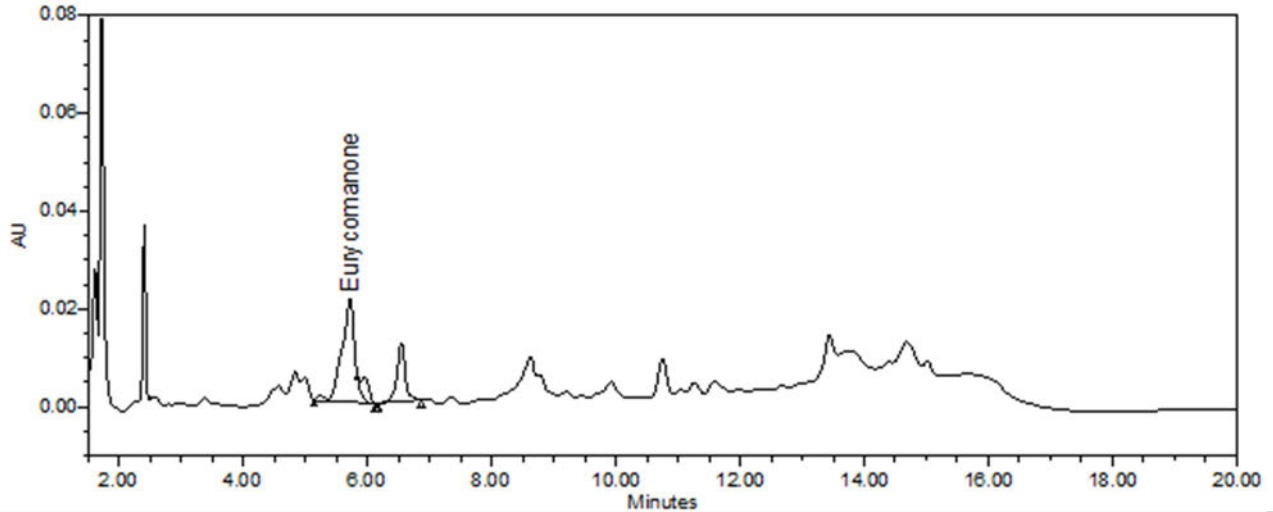


12661 HOOVER STREET GARDEN GROVE, CA 92841 | P. 714-754-4372 | F. 714-668-91

Report Issued To: Gorilla Mind, LLC
[Redacted]

Sample Name: Sigma
Capsule powder; Capsules in black finished
Description: product bottle
Lot #: 25010804
AL #: 25036FUM_3
Analysis ID: 253607
Received: 02/05/25

Determination of Eurycomanone Content by HPLC



Ret. Time (min)	Compound Name	Amount (%)	Amount (mg/serving)	Specification	Result
5.8	Eurycomanone	0.215	3.078	≥3 mg/serving	Pass

Chromatographic Conditions:

Method: ATM-815-0222
Column: AP392 Luna 5 µm C18(2) 100 A (150 x 4.6 mm)
Temperature: 40°C
Flow Rate: 1 mL/min
Injection Volume: 20 µL
UV Detection: 254 nm
Mobile Phase: 0.1% Formic Acid in Water
Acetonitrile
HPLC Instrument: Alliance_4

Sample Preparation:

Composited the contents of 10 capsules. Transferred approximately 300 mg of sample into a 50 mL volumetric flask. Added about 25 mL of methanol. Vortexed 30 seconds and sonicated 15 minutes at room temperature. Let cool and filled to volume with methanol. Filtered through 0.45 µm PTFE syringe filter into HPLC vial for analysis.

Report Summary:

Conclusion: This "Sigma" test sample contains 3 mg/2 capsules of Eurycomanone on the as-is basis.
Fill Weight: 716.239 mg/capsule; Serving size = 2 capsules.

OOS Reference: N/A
WI Reference: 04925 Eurycomanone

Analysis Date : 02/19/25
By: A Le

Analyzed



Digitally signed
by Latrece
Brown

Date:
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**Authorized By: Latrece Brown, Lead
Analytical Chemist**

The analytical method used has not been verified or validated for this product by Alkemist Labs. This report applies to the sample investigated and is not necessarily indicative of the quality or condition of apparently identical or similar products. This report is for the exclusive use of the party who requested the report and not for public dissemination or use by third parties, including for promotional purposes, without the prior written permission of Alkemist Labs. This report provides technical results for a specific sample and the report shall not be altered, modified, supplemented or abstracted in any manner. Any violation of these conditions renders the report and its results void. Pass/Fail decision is based on laboratory results as found.

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