



PERRY JOHNSON LABORATORY ACCREDITATION, INC.

Certificate of Accreditation

Perry Johnson Laboratory Accreditation, Inc. has assessed the Laboratory of:

Omnient Labs LLC

8370 S. Kyrene Road Tempe AZ 85284

(Hereinafter called the Organization) and hereby declares that Organization is accredited in accordance with the recognized International Standard:

ISO/IEC 17025:2017

This accreditation demonstrates technical competence for a defined scope and the operation of a laboratory quality management system (as outlined by the joint ISO-ILAC-IAF Communiqué dated April 2017):

Chemical Testing
(As detailed in the supplement)

Accreditation claims for such testing and/or calibration services shall only be made from addresses referenced within this certificate. This Accreditation is granted subject to the system rules governing the Accreditation referred to above, and the Organization hereby covenants with the Accreditation body's duty to observe and comply with the said rules.

For PJLA:

Tracy Szerszen
President

Initial Accreditation Date:

March 19, 2021

Issue Date:

April 27, 2023

Expiration Date:

May 31, 2025

Accreditation No.:

113861

Certificate No.:

L23-344

Perry Johnson Laboratory
Accreditation, Inc. (PJLA)
755 W. Big Beaver, Suite 1325
Troy, Michigan 48084

The validity of this certificate is maintained through ongoing assessments based on a continuous accreditation cycle. The validity of this certificate should be confirmed through the PJLA website: www.pjilabs.com



Certificate of Accreditation: Supplement

Omnient Labs LLC

8370 S. Kyrene Road, Tempe, AZ. 85284
 Contact Name: James Jursich Phone: 480-406-4034

Accreditation is granted to the facility to perform the following testing:

| FIELD OF TEST | ITEMS, MATERIALS OR PRODUCTS TESTED | SPECIFIC TESTS OR PROPERTIES MEASURED | SPECIFICATION, STANDARD METHOD OR TECHNIQUE USED | RANGE (WHERE APPROPRIATE) AND DETECTION LIMIT |
|-------------------------|--|--|---|---|
| Chemical ^F | Dietary Supplements – powders, capsules, tablets | Identity- Dietary supplement and botanical identity via High-Performance Thin Layer Chromatography (HPTLC) | Non-quantitative test HPTLC - AHPA or USP Botanical Reference Material | N/A – Qualitative method |
| | | Assay – Total Withanolides | HPLC – USP standard | Range: 0.01% to 100% (wt) LOD: 0.37 µg/g |
| | | Total Bacosides | UPLC – USP standard | Range: 0.01% to 100% (wt) LOD: 1.8 µg/g |
| | | Total Curcuminoids | HPLC – Sigma-Aldrich standard | Range: 0.001% to 100% LOD: 0.06 µg/g |
| | | Cordycepin | UPLC – Sigma-Aldrich standard | Range: 0.001% to 100% (wt) LOD: 0.001 µg/g |
| | | Total Silymarins | UPLC – Sigma-Aldrich standard | Range: 0.001% to 100% (wt) LOD: 2.0 µg/g |
| | | Eurycomanone | HPTLC – Sigma-Aldrich standard | Range: 0.1% to 100% (wt) LOD: 25 µg/g |
| | | Adrafinil | UPLC – Cayman Chemical Standard | Range: 0.1% to 100% (wt) LOD: 100 ng/g |
| | | Phenibut | UPLC – Cayman Chemical Standard | Range: 0.1% to 100% (wt) LOD: 100 ng/g |
| | | Phenylpiracetam | UPLC – Cayman Chemical Standard | Range: 0.1% to 100% (wt) LOD: 100 ng/g |
| | | Verbascoside and Echinacoside | UPLC – Sigma-Aldrich standard | Range: 0.1% to 100 (wt) LOD: 100 ng/g |
| | | Capsule mass | USP <2091> | As measured per USP guidelines |
| | | Tablet Disintegration | USP <701> | |
| | | Tablet Friability | USP <1216> | |
| | | Tablet hardness | USP <1217> | |
| Bulk and Tapped Density | USP <616> | | | |



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|-----------------------|--|--|--|--|
| Chemical ^F | Dietary Supplements – powders, capsules, tablets | Chemical Identity – FTIR Variable analytes | Non-quantitative test FTIR – Variable | N/A – Qualitative method |
| | | Total beta-D-glucans | UV-Vis – Megazyme Kit Standards 18% ± 3.0% | Range: 1% to 100% LOD: 1g/100g |
| | | Trace Analysis – Residual Solvents | USP <467> < 0.6% RSD average all classes | Range: 3 µg/g to 0.10 g/g LOD: Highly variable – to 0.37 µg/g for Class 1 solvents |

1. The presence of a superscript F means that the laboratory performs testing of the indicated parameter at its fixed location. Example: Outside Micrometer ^F would mean that the laboratory performs this testing at its fixed location.

