



PERRY JOHNSON LABORATORY ACCREDITATION, INC.

Certificate of Accreditation

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

***Biogen Laboratory Developments, LLC
Food, Supplement and R & D Lab
36740 E. Historic Columbia River Hwy, Corbett, OR 97019***

*(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):

***Biological (Microbiological) and 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:

April 09, 2024

Issue Date:

October 14, 2024

Expiration Date:

November 30, 2026

Accreditation No.:

75889

Certificate No.:

L24-782

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.pjllabs.com*



Certificate of Accreditation: Supplement

Biogen Laboratory Developments, LLC Food, Supplement and R & D Lab

36740 E. Historic Columbia River Hwy, Corbett, OR 97019
Contact Name: Mr. Nidal Kahl Phone: 503-695-6066

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

| FLEX CODE | FIELD OF TEST | ITEMS, MATERIALS, OR PRODUCTS TESTED | COMPONENT, CHARACTERISTIC, PARAMETER TESTED | SPECIFICATION OR STANDARD METHOD | TECHNOLOGY OR TECHNIQUE USED |
|-----------|--|--|--|--|---|
| F1, F2 | Biological ^F (Microbiological) | Food Products, Supplements, Nutraceuticals, Packaging Materials, and Environmental Samples | Aerobic Plate Count | AOAC 966.23 based on FDA BAM Online Ch.3; CMMEF 5 th Ed. | Culture – Pour Plate |
| F1, F2 | | | Yeast and Mold | SOP 220 based on FDA BAM Online Ch.18; CMMEF 5 th Ed | |
| F1, F2 | | | Anaerobic Plate Count | SOP 202 based on AOAC 966.23; FDA BAM Online Ch.3; CMMEF 5 th Ed. | |
| F1, F2 | | | Total Probiotic Count | CMMEF 5 th Ed. | |
| F1, F2 | | | Lactic Acid Bacteria | CMMEF 5 th Ed. | |
| F1, F2 | | | <i>Listeria monocytogenes</i> | AOAC 2019.11 | GENE-UP [®] PCR |
| F1, F2 | | | <i>E. coli</i> O157:H7 | AOAC 2019.03 | |
| F1, F2 | | | EHEC Series: STEC – EAE & STX STEC - top 6 | AOAC PTM 121806 | GENE-UP [®] PCR & Vidas [®] Immunoassay |
| F1, F2 | | | <i>Salmonella sp.</i> | AOAC 2020.02 PEPRL 0007.10 | GENE-UP [®] PCR |
| F1, F2 | | | <i>Salmonella sp.</i> | AOAC 2013.01 | Vidas [®] UP Immunoassay |
| F1, F2 | | | <i>Listeria sp.</i> | AOAC 2013.10 | |
| F1, F2 | | | <i>Listeria monocytogenes</i> | AOAC 2013.11 | Vidas [®] Xpress Immunoassay |
| F1, F2 | | | <i>Listeria monocytogenes</i> | AOAC 2004.02 | Vidas [®] LMO2 Immunoassay |
| F1, F2 | | | <i>E. coli</i> O157:H7 | AOAC PTM 060903 | Vidas [®] UP Immunoassay |
| F1, F2 | | | <i>Salmonella sp.</i> | AOAC 2013.02 | BAX [®] System PCR |
| F1, F2 | | | <i>Listeria sp.</i> | AOAC PTM 081401 AOAC PTM 050903 | |
| F1, F2 | | | <i>Listeria monocytogenes</i> | AOAC PTM 121402 AOAC PTM 080901 | |
| F1, F2 | | | HEC Series: STEC – EAE & STX STEC – Panel 1 & 2 | AOAC PTM 091301 USDA FSIS MLG 5B.05 | |
| F1, F2 | | | Campylobacter | AOAC PTM 040702 USDA FSIS MLG 41.04,41A.00 | |
| F1, F2 | | | <i>Vibrio cholerae/parahaemolyticus/vulnificus</i> | AOAC PTM 050902 | |



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|------------|--|--|---|--|--------------------------------|----------------------------------|-----------------------------------|
| F1, F2 | Biological ^F (Microbiological) | Food Products, Supplements, Nutraceuticals, Packaging Materials, and Environmental Samples | Salmonella sp. | AOAC RI 080601 | RapidChek Immunoassay | | |
| F1, F2 | | | Listeria sp. | AOAC RI 020401 | | | |
| F1, F2 | | | E. coli O157:H7 | AOAC RI 070801 | | | |
| F1, F2 | Biological ^F (Microbiological) | Food Products, Supplements, Packaging Materials, and Environmental Samples | Aerobic Plate Count (Rapid) | AOAC 2015.13 | Petrifilm | | |
| F1, F2 | | | Total Coliform | AOAC 991.14 | | | |
| F1, F2 | | | Lactic Acid Bacteria | AOAC RI 041701 | | | |
| F1, F2 | | | Total E. coli (Generic) | AOAC 991.14 | | | |
| F1, F2 | | | Staphylococcus aureus | AOAC 2003.07 | | | |
| F1, F2 | | | Total Enterobacteriaceae | AOAC 2003.01 | | | |
| F1, F2 | | | Yeast and Mold(Rapid) | AOAC 2014.05 | | | |
| F1, F2 | | | Drinking Water | Total Coliform | | SM 9223B | Colilert |
| F1, F2 | | | | Total E. coli (Generic) | | SM 9223B | |
| F1, F2, F4 | | | Biological ^{FO} (Microbiological) | Food Products, Supplements, Nutraceuticals, Packaging Materials, and Environmental Samples | | Microbial Environmental Sampling | CMMEF 5 th Ed. SOP 300 |
| F1, F2 | Biological ^F (Microbiological) | Food Products, Supplements, Nutraceuticals, Packaging Materials, and Environmental Samples | Listeria sp. | AOAC 2019.10 | GENE-UP [®] PCR | | |
| F1, F2 | | | Cronobacter sp. | AOAC 2019.01 | | | |
| F1, F2 | | | Campylobacter | AOAC PTM 051201 USDA FSIS MLG 41.01 | Vidas [®] Immunoassay | | |
| F1, F2 | | | E. coli O157:H7 | AOAC PTM 102003 AOAC PTM 031002 USDA FSIS MLG 5.09/5A.04 | BAX [®] System PCR | | |
| F1, F2 | | | Listeria monocytogenes | AOAC RI 011805 | RapidChek Immunoassay | | |
| F1, F4 | | Food: Fruit, Vegetables, Water, Surfaces | Norovirus GI-GII, Hepatitis A Virus | SOP 352 based on ISO 15216 | RT-qPCR | | |
| F1, F4 | Chemical ^F | Fruit, Plant Tissue, Supplements, Nutraceuticals | Pesticide Analysis | AOAC 2007.01 modified SOP 313 | LC-MS/MS | | |
| F1, F4 | | | | GC-MS/MS | | | |
| F1, F4 | | | Herbicide Analysis | AOAC 2007.01 modified SOP 313 | LC-MS/MS | | |
| F1, F4 | | | Cannabinoid Profile | SOP 620 | HPLC | | |



Certificate of Accreditation: Supplement

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|-----------|--|--------------------------------------|---|-------------------------------------|---------------------------------|
| F1, F4 | Chemical ^F | Cannabis, Oils, Distillates | Pesticide Analysis | SOP 630 AOAC 2007.01 modified | LC-MS/MS |
| F1, F4 | | | | SOP 630 AOAC 2007.01 modified | GC-MS/MS |
| F1, F2 | Food Products, Supplements, Nutraceuticals, Packaging Materials, and Environmental Samples | | Gliadin R5 / Gluten | AOAC RI 061201 | ELISA |
| F1, F4 | | | Sesame Allergen | ELISA SOP 262 | |
| F1, F4 | | | Egg Allergen | ELISA SOP 251 | |
| F1, F4 | | | Milk Allergen | ELISA SOP 252 | |
| F1, F4 | | | Soy Allergen | ELISA SOP 253 | |
| F1, F2 | | | Peanut Allergen | AOAC RI 030403 | |
| F1, F4 | | | Crustacea Allergen | ELISA SOP 255 | |
| F1, F4 | | | Tree Nut Allergen | ELISA SOP 259 | |
| F1, F2 | | | Mycotoxins: Aflatoxin B1, B2, G1, G2 | AOAC RI 050901 | ELISA |
| F1, F4 | | | Mycotoxin: Aflatoxin M1 | ELISA SOP 271 | |
| F1, F2 | | | Mycotoxin: DON / Vomitoxin | AOAC RI 090901 | |
| F1, F4 | | | Mycotoxin: Ochratoxin | SOP 273 | |
| F1, F2 | | | Water Activity | AOAC 978.18 | Aw Meter |
| F1, F2 | | | Moisture | AOAC 925.10 | Gravimetric |
| F1, F2 | | | Solids | AOAC 925.10 | |
| F1, F2 | | | pH | AOAC 981.12; AOAC 960.19 | pH Meter |
| F1, F4 | | | Food Products, Supplements, Nutraceuticals | Patulin | AOAC 995.10 modified SOP 306 |



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1. The presence of a superscript F means that the laboratory performs testing of the indicated parameter at its fixed location.
2. The presence of a superscript O means that the laboratory performs testing of the indicated parameter onsite at customer locations.
3. Flex Code:
 - F0-Fixed scope item. No deviations allowed to the line item as identified, except for updating to the most recent version of an accredited standard method after verification
 - F1-Laboratory has the capability to test a new item, material, matrix, or product similar in composition to item, material, matrix, or product identified on the scope
 - F2-Laboratory has the capability to introduce the newest revision of an accredited authoritative standard method (with no modifications) identified on the scope
 - F3-Laboratory has the capability to introduce a parameter/component/analyte to an accredited test method identified on the scope
 - F4-Laboratory has the capability to introduce a new revision of an accredited non-standard method using the same technology or technique identified on the scope
 - F5-Laboratory has the capability to introduce a validated method that is equivalent to an accredited method (using same technology or technique) identified on the scope