PERRY JOHNSON LABORATORY ACCREDITATION, INC.

Certificate of Accreditation

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

Consumer Product Testing Company, Inc.

70 New Dutch Lane, Fairfield, NJ 07004

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

ISO/IEC 17025:2005

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 January 2009):

Biological 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/Operations Manager

Initial Accreditation Date:	*Issue Date:*	*Expiration Date:*
September 29, 2016	September 29, 2016	September 29, 2018

Accreditation No.:	*Certificate No.:*
80071	L16-407

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.pjlabs.com

Certificate of Accreditation: Supplement

Consumer Product Testing Company, Inc.

70 New Dutch Lane, Fairfield, NJ 07004
Contact: William Neumann Phone: 973-808-7111

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
Biological[F]	Ingredients, Drug Products, Cosmetic and Consumer Products, Dietary Supplements and Medical Devices	Microbial Content	Harmonized USP 61 & 62 and EP 2.6.12, 2.6.13	Presence/Absence or CFU/g or mL Varies with analyte
		Sterility	USP <71> EP 2.6.1 AAMI/ISO 11737-2:2009	
		Bioburden	AAMI 11737-1: 2006	
		LAL Endotoxin	USP <85> EP 2.6.14	
		Preservative Effectiveness	USP <51> EP 5.1.3 CTFA	
		Cosmetic Products Preservative Effectiveness	ISO 11930:2012	
		Water Testing	CPTC010020	
		Ames Mutagenicity (Bacterial Reverse Mutation Assay)	OECD 471	
Chemical[F]	Ingredients, Drug Products, Dietary Supplements and Consumer Products	Residual Ethylene Oxide and Dioxane	USP <228> Ethylene Oxide and Dioxane	N/A
			Method 1	ETO 10 ppm Dioxane 10 ppm
			Method 2	ETO 1 ppm Dioxane 10 ppm
		Determination of residual solvents	USP <467> Residual Solvents GC	Varies with analyte ppm levels
		Potency and Impurity Assays	USP<621> Chromatography HPLC and GC	Typically % or mg quantities Varies with analyte
	Children's Toys and Childcare Articles	Determination of Phthalates	CPSC-CH-C1001-09.3	Typically % or mg quantities

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.

2. "Accreditation is granted through technology based flexible scope criteria. Additional methods other than listed above may fall under the accreditation of the laboratory. A complete listing of method capabilities can be derived from the laboratory upon request".