

Natural-Immunogenics Corporation
Standard Operating Procedure

Department: Regulatory & Quality Control

SOP MFG-33 Current Revision

Homeopathic Gel

Certificate of Analysis

Regulatory Compliance: International regulatory standards for the manufacture of Homeopathic Products and OTC Pharmaceuticals require the Issuance of a COA for each bulk batch of product to confirm that specifications are met in terms of product identity, purity, strength and composition. Complete quality & purity testing were performed as delineated in the MBR and results are recorded in the batch record. This batch is hereby certified to have been produced in accordance with the Formulary, Quality controls, Specifications and Characteristics set forth by NIC proprietary formulations. There have been no revisions or changes, which have occurred during the production of this batch except as noted herein.


Bulk Batch No: G001120419 **Filled Lot:** HM029T **Expiry:** 10/2022

ANALYTICAL DATA	PRODUCT SPECIFICATIONS	TEST RESULTS	TEST METHOD USED
VISCOSITY: SP # 6 @ 25°C	40,000 – 55,000cPs	52,017	SOP LAB-54
pH	4.5 to 5.5	5.3	SOP LAB-02
COLOR	Clear to Slightly Tinted	Clear	VISUAL
ODOR	Odorless	Odorless	OLFACTORY

MICROBIAL PURITY TESTING

ANALYTICAL DATA	PRODUCT SPECIFICATIONS	TEST RESULTS	TEST METHOD USED
TOTAL AEROBIC COUNT	<100 CFU/mL	Pass	EPA APPROVED TEST METHOD; SOP LAB-91
TOTAL MOLD AND YEAST	<10 CFU/mL	Pass	EPA APPROVED TEST METHOD; SOP LAB-91

DATA TRANSFERRED FROM ORIGINAL COA BY:

PRINT NAME	TITLE	SIGNATURE & DATE
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