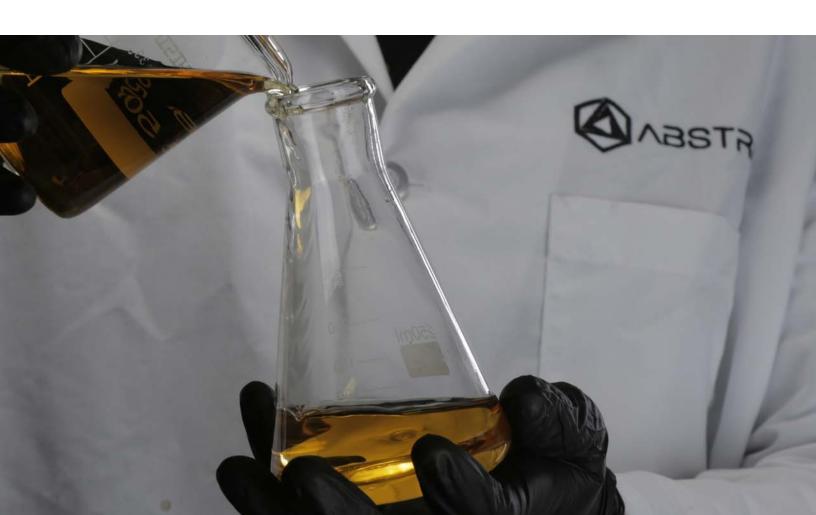


Compliance Packet

Rev.5 | 12.1.2021



ABOUT US

Abstrax is a collection of scientists, cultivators, entrepreneurs, and passionate advocates working together to push the limits of terpene product formulations. More than that, we are a family of explorers pioneering the frontiers of sensory design and terpene manufacturing for the benefit of all.

MISSION

To unravel the secrets of cannabis terpenes and aroma compounds. To discover new compounds and use them to empower the community. To set the standards and best practices that allow brands to differentiate their products. To change the landscape of transportation and provide consistent experiences for brand owners. To be as innovative as we are transparent. To take what nature has given us and perfect it with science.

OUR VISION

The creation of a modern, global industry built around terpenes and their applications. Perpetual innovation and strategic collaboration. Engage, inspire, and empower our clients to become successful pioneers of sensorial experiences.

ABOUT OUR TERPENES

Abstrax knows what it takes to win extract awards. Our adherence to quality begins with the absolute best starting materials. Abstrax terpenes and terpenoids are isolated from natural source materials found all around the world. They are distilled multiple times to achieve the absolute highest purity and do not contain CBD, THC or any other illegal substances.

Our terpenes are 100% free of fillers like PG, VG, MCT and coconut oil. No added fillers equates to low dose and high potency in your product formulations. Additionally, our terpenes and terpene strain profiles are derived from natural ingredients, solvent-free, GMO-free, filler-free and food grade.

Thus, we will not offer them for sale until we know they are a perfect fit. Every strain profile is originally based upon several lab results of the same strain that are continually analyzed and further refined through a series of proprietary testing protocols.

These tests ensure that the flavor matches that of the unique flower as closely as humanly possible. We offer proven strain profiles for mixing, isolates for experimenting with new flavors, or custom solutions for those seeking unique formulations.

TABLE OF CONTENTS

SELF-AUDIT FORM	1
ISO 9001: 2015	9
HACCP CERTIFICATE	10
FOOD GRADE STATEMENT	11
PESTICIDE AND HEAVY METAL DECLARATION	12
NO RISK OF SALMONELLA DECLARATION	13
NON-GMO DECLARATION	14
NATURAL FLAVOR STATEMENT	15
COUNTRY OF ORIGIN STATEMENT	16
TSE BSE STATEMENT	17
BEST BY STATEMENT	18
DIETARY PREFERENCES & RESTRICTIONS STATEMENT	19
PHARMACEUTICAL EXCIPIENTS STATEMENT	20
VITAMIN E ACETATE STATEMENT	21
PHYTOL STATEMENT	21
VAPE CARTRIDGE INGREDIENTS STATEMENT	22
KOSHER CERTIFICATE	24
QUALITY STATEMENT	25
QUALITY POLICY	25
TERPENE DISTILLATION PROCESS FLOW DIAGRAM	26
TERPENE BLENDING PROCESS FLOW DIAGRAM	27
HYGIENE POLICY	28
ALLERGEN / SENSITIVE AGENTS IDENTIFICATION SHEET	29
GAS CHROMATOGRAPHY ANALYSIS COMPOUND TEST	30
FREQUENTLY ASKED QUESTIONS	32
CERTIFICATE OF ANALYSIS	34
SAFETY DATA SHEET	36
PRODUCT DATA SHEFT	41

Section 1: General Information

Company Name	Abstrax Tech
Business Type	Manufacturer
Product Description or Service	Terpene Hydrocarbons
Mailing Address	2661 Dow Ave. Tustin, CA 92780
Phone Number	(562) 294 - 5805
Email	info@abstraxtech.com
Is the company private or public?	Private
Number of years in business?	3 Years
Number of Personnel?	40 Employees
Is your company currently increasing or decreasing employee levels?	Increasing
What is the square footage of the facility?	10,400 sq ft.
Do you operate locally, nationally or globally?	Locally
Number of personnel in Production?	22
How many shifts?	Single 8-hr shift
Number of personnel in Administration?	10
Number of personnel in Operations?	5
QC Contact	Name, Title: Donald Remmel, QC/Inventory Analyst
Number of Personnel in QC?	3
Is the QC department independent of production?	Yes
Does your organization have a business license?	Yes - City of Tustin Business License, Tax Certificate: 257643008-00002
Is the facility FDA registered?	Yes
Operating in accordance with GMP practices?	Yes - Incorporated in HACCP program
Is your organization HACCP certified?	Yes - HACCP certification
Is your organization ISO9001:2015 certified?	Yes
Is your organization FEMA GRAS registered?	Yes
Is your organization EPA registered?	Yes
Is your organization hazmat shipping certified?	Yes
Are you a licensed food manufacturer?	No

Section 2: Quality Systems

	Yes	No	N/A	Comments
Do you have a quality system?	$\langle \hspace{-0.5em} \hspace{-0.5em} \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	A Table of Contents is attached.
Do you have a quality policy statement?	$\langle \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \!$	\bigcirc	\bigcirc	Listed on page 11.
Is the policy communicated to all staff and understood?	$\langle \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	
Is the quality manual kept current and available to employees, auditors, or customers?	$\langle \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \!$	\bigcirc	\bigcirc	
Is there a process in place to cover for absence of key staff?	$\langle \hspace{-1em} \hspace{-1em} \hspace{-1em} \hspace{-1em} \hspace{-1em} \rangle$	\bigcirc	\bigcirc	
Do you have quality objectives for production department?	$\langle \hspace{-0.5em} $	\bigcirc	\bigcirc	
Is the Quality department independent of operations?	$\langle \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	
Are there formal written procedures for tests performed?	$\langle \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	
Do you have a customer complaint handling procedure?	$\langle \hspace{-1em} \hspace{-1em} \rangle$	\bigcirc	\bigcirc	
Is the Management review of the quality system on a clearly defined schedule and documented?	$\langle \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \!$	\bigcirc	\bigcirc	
Do you have an internal audit program?	$\langle \hspace{-0.5em} \hspace{-0.5em} \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	
Are there internal audits scheduled and carried out?	$\langle \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	
Do you have standard operating procedures?	$\langle \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	
Is there a document control system in place?	$\langle \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	
Are Certificate of Analysis (Conformance) issued with your products on every order?	$\langle \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \!$	\bigcirc	\bigcirc	
Are nonconformance's investigated?	$\langle \hspace{-0.5em} \hspace{-0.5em} \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	
Do you have instruments that require calibration?	$\langle \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	
Are records kept of all control results?	$\langle \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	
If yes for how long do you keep these records?	$\langle \hspace{-0.5em} \hspace{-0.5em} \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	7 Years
Is there a record keeping system along with a revision control system for quality documentation?	$\langle \hspace{-0.5em} \hspace{-0.5em} \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	
Would you provide prior notification for changing suppliers and manufacturing site? Change notification policy?	$\langle \hspace{-1em} \hspace{-1em} \rangle$	\bigcirc	\bigcirc	
Have you been audited by other companies?	$\langle \hspace{-1em} \hspace{-1em} \rangle$	\bigcirc	\bigcirc	
Is there a documented process and performance metric used to measure the level of customer satisfaction?	$\langle \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \!$	\bigcirc	\bigcirc	
Do you have a method for testing finished product?	$\langle \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \!$	\bigcirc	\bigcirc	
Are samples from each lot retained? If so, how long?	$\langle \overline{\boldsymbol{\mathcal{E}}} \rangle$		\bigcirc	Retained samples are kept in a temperature – controlled environment and are retained for up to 2 years.

Section 2: Quality Systems (cont.)

	Yes	No	N/A	Comments
Does the company operate a formal system of training, including new hire training with records maintained and reviewed periodically?	$\langle \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \!$	\bigcirc	\bigcirc	Job-specific, is performed as per HACCP requirements.
Is there a documented recall plan in place?	$\langle \hspace{-1em} \hspace{-1em} \hspace{-1em} \hspace{-1em} \rangle$	\bigcirc	\bigcirc	
Do you conduct a mock recall? If so, how often?	$\langle \hspace{-0.5em} \hspace{-0.5em} \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	Annually
Do you have a method to notify customers in the event of a recall?	$\langle \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	
Do you document the recall event?	$\langle \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	
Do you have a formalized root cause, corrective and preventive procedure for non-conformances?	$\langle \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \!$	\bigcirc	\bigcirc	
Is there an established and documented internal audit/self-evaluation program that promotes continuous improvement?	$\langle \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	
Has the competency of internal auditors been assessed and do you ensure they do not audit areas which they are responsible for?	$\langle \hspace{-1em} \hspace{-1em} \rangle$	\bigcirc	\bigcirc	
Has a schedule been developed outlining the scope of the audit, audit frequency, and corrective action monitoring/follow-up?	$\langle \hspace{-1em} \hspace{-1em} \rangle$	\bigcirc	\bigcirc	
Is there record control of audit and corrective action documentation along with verification of compliance? Has this documentation been reviewed by Quality Management?	$\langle \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \!$	\bigcirc	\bigcirc	

Section 3: Facilities and Equipment

	Yes	No	N/A	Comments
Do you use a facility safety checklist?	$\langle \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	
Do you inspect the facility? If so, how often?	$\langle \hspace{-0.5em} \hspace{-0.5em} \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	Monthly
Is there adequate security to prevent entry by unauthorized personnel?	$\langle \hspace{-1em} \hspace{-1em} \hspace{-1em} \hspace{-1em} \rangle$	\bigcirc	\bigcirc	
Do you have a Preventative Maintenance Program?	$\langle \hspace{-1em} \hspace{-1em} \rangle$	\bigcirc	\bigcirc	
Is all key production equipment included in the Preventative Maintenance Program?	$\langle \!$	\bigcirc	\bigcirc	
Is there a method to prevent cross-contamination?	\bigcirc	\bigcirc	$\langle \hspace{-0.5em} \hspace{-0.5em} \hspace{-0.5em} \hspace{-0.5em} \rangle$	We have no allergens within the facility
Are facilities maintained and in good state of repair?	$\langle \hspace{-1em} \hspace{-1em} \hspace{-1em} \hspace{-1em} \rangle$	\bigcirc	\bigcirc	
Do you have a cleaning/sanitation program?	$\langle \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \!$	\bigcirc	\bigcirc	
Do you use clean water that is contamination-free?	$\langle \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \!$	\bigcirc	\bigcirc	Water is not used in production, only in cleaning of the facilities and glassware.
Do you segregate clean/dirty equipment?	$\langle \hspace{-1em} \hspace{-1em} \hspace{-1em} \hspace{-1em} \rangle$	\bigcirc	\bigcirc	
Do you have a process for handling/removing waste?	$\langle \hspace{-0.5em} \hspace{-0.5em} \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	
Do you use covered waste containers?	$\langle \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	
Do you use pallets for your facility? If so, what kind?	$\langle \hspace{-0.5em} \hspace{-0.5em} \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	Wooden/Plastic
Are all pertinent production equipment included in a preventive maintenance program?	$\langle \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \!$	\bigcirc	\bigcirc	
Do you have a calibration system and is it traceable to a known standard?	$\langle \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc		Testing and measuring equipment calibrated by 3rd party.
Is there adequate space, lighting and appropriate racks to prevent damage or mishandling of materials?	$\langle \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \!$	\bigcirc	\bigcirc	

Section 4: Safety, Health and Environment

	Yes	No	N/A	Comments
Do you have an operational management system for Safety, Health and Environment?	$\langle \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	
System is based on HACCP?	$\langle \overline{\mathbf{V}} \rangle$	\bigcirc	\bigcirc	
Do you have a dedicated team for safety and health?	$\langle \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	
Do you have a safety team that regularly updates a Hazard Analysis that identifies all hazards associated with your facility?	$\langle \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	
Are all identified hazards controlled by your facility?	$\langle \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	
Have employees been trained in handling the hazardous waste produced by your manufacturing process?	$\langle \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	
Is your hazardous waste removed and disposed of by a 3rd party?	$\langle \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	Environmental Logistics
Is the process of having your hazardous waste materials properly removed within the required time frame?	$\langle \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \!$	\bigcirc	\bigcirc	Yes, employees have been trained to contact 3rd party prior to 180 days.
Do you have programs to control and reduce waste and control the environmental impact?	$\langle \hspace{-1em} \hspace{-1em} \rangle$	\bigcirc	\bigcirc	

Section 5: Sanitation and Hygiene

	Yes	No	N/A	Comments
Do you have sanitation control SOPs?	$\langle \hspace{-0.5em} \hspace{-0.5em} \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	
Do you have cleaning schedules?	$\langle \hspace{-1em} \hspace{-1em} \hspace{-1em} \hspace{-1em} \hspace{-1em} \rangle$	\bigcirc	\bigcirc	
Do you keep records of these?	$\langle \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	Glassware and contact surfaces are cleaned/sanitized.
Are cleaning schedules reviewed/verified?	$\langle \hspace{-0.5em} \hspace{-0.5em} \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	
Does the company require that employees report to work in good health, clean and dressed in clean attire?	$\langle \hspace{-0.6em} \hspace{-0.6em} \rangle$	\bigcirc	\bigcirc	
Are chemicals segregated from other ingredients, correctly labeled, and stored?	$\langle \hspace{-0.5em} \hspace{-0.5em} \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	Chemicals stored away from Raw Materials and Finished Goods.
Are employees trained on hygiene?	$\langle \hspace{-1em} \hspace{-1em} \hspace{-1em} \hspace{-1em} \hspace{-1em} \rangle$	\bigcirc	\bigcirc	
Is smoking permitted in designated areas only?	$\langle \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	Designated areas are outside of the building.
Is eating and drinking permitted in designated areas only?	$\langle \hspace{-1em} \hspace{-1em} \rangle$	\bigcirc	\bigcirc	
Are sick personnel and visitors prevented from entering production areas?	$\langle \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \!$	\bigcirc	\bigcirc	Sick workers are not allowed into production.
Are coverings to minor injuries required to be covered by a bandage?	$\langle \overline{\boldsymbol{V}} \rangle$	\bigcirc	\bigcirc	Bandage and single use gloves are required for injured hands.
Are all production personnel required to wear hair/beard nets for product protection?	$\langle \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \!$	\bigcirc	\bigcirc	PPE signs are posted at each production entrance.
Are lab coats provided?	$\langle \hspace{-0.5em} $	\bigcirc	\bigcirc	
Are handwashing facilities provided?	$\langle \hspace{-1em} \hspace{-1em} \rangle$	\bigcirc	\bigcirc	
Are handwashing signs visible and legible?	$\langle \hspace{-0.5em} \hspace{-0.5em} \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	
Are restrooms separate from production areas?	$\langle \hspace{-0.5em} \hspace{-0.5em} \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	
Are lockers provided for employees?	$\langle \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \!$	\bigcirc	\bigcirc	
Is hand soap provided?	$\langle \hspace{-0.5em} \hspace{-0.5em} \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	
Are paper towels available for drying hands?	$\langle \hspace{-0.5em} \hspace{-0.5em} \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	

Section 6: Pest Control

	Yes	No	N/A	Comments
Do you have a pest control program?	$\langle \hspace{-1em} \hspace{-1em} \rangle$	\bigcirc	\bigcirc	Western Exterminator
Do trained personnel carry out pest control?	$\langle \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \!$	\bigcirc	\bigcirc	
Are all pest control stations identified?	$\langle \hspace{-1em} \hspace{-1em} \rangle$	\bigcirc	\bigcirc	
Are corrective actions documented?	$\langle \hspace{-1em} \hspace{-1em} \hspace{-1em} \hspace{-1em} \rangle$	\bigcirc	\bigcirc	
Are there enough pest stations for the facility?	$\langle \hspace{-1em} \hspace{-1em} \hspace{-1em} \hspace{-1em} \rangle$	\bigcirc	\bigcirc	
Are windows and doors to production areas adequate to prevent ingress of pests?	$\langle \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \!$	\bigcirc	\bigcirc	

SELF-AUDIT FORM

Section 7: Packaging, Labeling and Shipping

	Yes	No	N/A	Comments
Is each container labeled with the lot/batch number?	$\langle \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	
Do you keep records of all shipments to customers, including batch number and quantity?	$\langle \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \!$	\bigcirc	\bigcirc	
Are raw materials and finished products stored in clean, dry, and wellventilated spaces and protected from contamination?	$\langle \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \!$	\bigcirc	\bigcirc	
Do you have a regular carrier for your goods?	$\langle \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \!$	\bigcirc	\bigcirc	UPS/FedEx
What is the name of the name of the company?	$\langle \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \!$	\bigcirc	\bigcirc	UPS/FedEx
Who is the carrier agent?	$\langle \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \!$	\bigcirc	\bigcirc	UPS/FedEx
Is packaging stored in a safe area?	$\langle \hspace{-1em} \hspace{-1em} \rangle$	\bigcirc	\bigcirc	
Does your transport system make use of a tracking report?	$\langle \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \!$	\bigcirc	\bigcirc	
Is traceability for packing ensured?	$\langle \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \!$	\bigcirc	\bigcirc	
Is the packaging tamper evident?	$\langle \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \!$	\bigcirc	\bigcirc	

SELF-AUDIT FORM

Section 7: Packaging, Labeling and Shipping (cont.)

	Yes	No	N/A	Comments
Do you have an internal laboratory?	$\langle \hspace{-1em} \hspace{-1em} \rangle$	\bigcirc	\bigcirc	
Is an outside laboratory used for any testing?	$\langle \overline{\mathbf{V}} \rangle$	\bigcirc	\bigcirc	
What types of testing is contracted out?	$\langle \hspace{-1em} \hspace{-1em} \rangle$	\bigcirc	\bigcirc	Heavy Metals and Pesticide (Random) Testing
Have you qualified/evaluated these contract laboratories?	$\langle \hspace{-1em} \hspace{-1em} \rangle$	\bigcirc	\bigcirc	
Do you use skip lot testing?	\bigcirc	$\langle \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	Every lot is tested.
Do you have procedures for minimizing packaging and labelling errors?	$\langle \hspace{-1em} \hspace{-1em} \hspace{-1em} \hspace{-1em} \rangle$	\bigcirc	\bigcirc	

SELF-AUDIT FORM

Section 8: Item/Lot Number Assignment

	Yes	No	N/A	Comments
Do you have a specific method for issuing lot/batch numbers?	$\langle \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	Lot numbers are automatically issued by our ERP
Is there a system of product batch/lot control for identification and traceability purposes?	$\langle \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	
Does the Quality Department inspect or verify each batch/lot for conformance prior to release and shipment to customers?	$\langle \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	
Is there a product recall control system in place which ensures that any materials shipped can be traced and recalled?	$\langle \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	
Are non-conforming materials requiring re-work segregated and stored in a location protected from the environment?	$\langle \hspace{-0.5em} \hspace{-0.5em} \rangle$	\bigcirc	\bigcirc	
Is there a system of issuing a compliance statement or certification of analysis to customers ensuring the material meets specifications?	$\langle \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \! \!$	\bigcirc	\bigcirc	

The following documentation is provided on the website: Product details, SDS, Specifications and COA's. To obtain additional specific information regarding please e-mail us at info@abstraxtech.com.



Certificate of Registration

Perry Johnson Registrars, Inc., has audited the Quality Management System of:

Abstrax Tech, Inc.
2661 Dow Avenue, Tustin, CA 92780 United States

(Hereinafter called the Organization) and hereby declares that Organization is in conformance with:

ISO 9001:2015

This Registration is in respect to the following scope:

Design, Manufacture, Distribution and Research of Terpene Blends

This Registration is granted subject to the system rules governing the Registration referred to above, and the Organization hereby covenants with the Assessment body duty to observe and comply with the said rules.







Terry Boboige, President

Perry Johnson Registrars, Inc. (PJR) 755 West Big Beaver Road, Suite 1340 Troy, Michigan 48084 (248) 358-3388

The use of the UKAS accreditation symbol is in respect to the activities covered by the Accreditation Certificate Number 0105.

The validity of this certificate is dependent upon ongoing surveillance.

Effective Date: Revision Date:

Certificate No.:

March 2, 2020

May 24, 2021

Expiration Date: March 1, 2023

C2020-00829-R1





FOOD GRADE STATEMENT

This is to verify that all flavor products produced by Abstrax Tech Inc., are of food grade quality. The materials used are allowed for use as a flavor ingredient intended for human consumption, or as an indirect additive according to the United States Food & Drug Administration 21 CFR and complies with European 88/388/EEC and 1334/2008/CE Flavorings directive. The materials used are manufactured in accordance with Good Manufacturing Practices (GMPs).

Donald Remmel

QC/Inventory Analyst

Donald Hunn



PESTICIDE AND HEAVY METAL DECLARATION

We hereby confirm that the ingredients used in the manufacture of our flavor and fragrance products meet the E.U. regulations as outlined in 1881/2006/EC in regard to contaminants prohibited in food grade raw material ingredients. These ingredients also comply with the E.U. regulation 396/2005/EC, regarding the maximum residue levels of pesticides in food grade raw material ingredients of plant and animal origin. The materials used are manufactured in accordance with Good Manufacturing Practices. (GMP's).

Donald Remmel

Yould Theners

QC/Inventory Analyst



March 30, 2021

Declaration of No Risk of Salmonella

At Abstrax Tech Inc. our Terpene blends are created with the finest ingredients and manufactured under safe monitored conditions. We are HACCP certified and there is no risk of any products containing Salmonella.

Donald Remmel

Donald Henry

QC/Inventory Analyst



April 23, 2021

NON-GMO DECLARATION

This letter is to certify products offered and sold by Abstrax Tech., Inc. are not derived from or produced using GMO's or their derivatives, and that all reasonable steps have been taken to avoid contamination from GMO or their derivatives.

Donald Remmel

QC/Inventory Analyst

Donald Henry



NATURAL FLAVOR STATEMENT

All Abstrax Tech., Inc. Natural blend products are blends that include natural terpenes and may include natural flavorings. The term Natural is per the definition contained in CFR§ 101.22(a)(1)-(3).

Donald Remmel

QC/Inventory Analyst

Donald Theren



COUNTRY OF ORIGIN STATEMENT

This letter is to certify that all products sold by Abstrax Tech Inc., at the Tustin, California facility are wholly manufactured in the United States of America. All products and packaging meet FDA requirements.

Donald Remmel

QC/Inventory Analyst

Donald Thenens



STATEMENT ON TRANSMISSIBLE SPONGIFORM ENCEPHALOPATHY (TSE) and BOVINE SPONGIFORM ENCEPHALOPATHY (BSE)

At Abstrax Tech Inc. our products do not contain any raw materials produced from or substances derived of animal origin. Moreover, these products do not contain, and are not derived from, specified risk material as defined in Commission Decision 97/534/EC or mechanically recovered meat obtained from the vertebral column of bovine, ovine or caprine animals.

The manufacturing process does not use any ingredient of animal origin nor do our products come in contact with animal products during storage and transportation.

Therefore, to the best of our knowledge, all of our products are free from Transmissible Spongiform Encephalopathy (TSE) and Bovine Spongiform Encephalopathy (BSE).

Donald Remmel

Sould Thurst

Quality Control Inventory Analyst



2661 Dow Avenue, Tustin, CA 92780

April 26, 2021

STORAGE CONDITIONS AND IMPACT ON SHELF LIFE Terpene Flavor Blends

Shelf Life

The shelf life statements on Abstrax product labels and supplemental product documentation, reflect the product's "Fitness for Use." The term "Fitness for Use" indicates acceptable terpene flavoring characteristics, as opposed to food safety concerns that would render the product unusable. Natural terpene flavors are somewhat dynamic. They can undergo shifts in flavor, color and /or pH as the products age. Some products are more susceptible to shifts than other products. Storage conditions will exacerbate or delay these shifts. Therefore, Abstrax recommends keeping the terpene flavors under the recommended storage conditions. Conditions that can negatively affect the shift in shelf-life include exposure to heat, light, high moisture, or freezing depending on the product.

Open Containers

The shelf life of opened natural terpene flavors will be dependent on the age of the product and on storage conditions. It is advantageous to use the natural terpene flavors expeditiously after opening. If the natural terpene flavor must be carried over after use, the container should be tightly sealed, and be stored under the optimal conditions. Partial containers should be used as soon as possible for best results. Opened containers should be usable as long as the product remains characteristic to target specifications.

Shelf Life Extensions

Abstrax strongly encourages customers to use our natural terpene flavors within the stated shelf life. The customer should be aware that the flavor, color and/or pH characteristics of natural terpene flavors will shift with age. Some products may function well after the shelf life date. Abstrax will not warrant any product's performance after the expiration of shelf life. Any attempt to use expired product will be at the sole discretion of the customer and should be based on the customer's testing in their products and formulations.

Donald Remmel

Yould Though

QC/Inventory Analyst



15550 Rockfield Blvd., Suite B120, Irvine, California 92618

February 2, 2021

Dietary Preferences & Restrictions

At Abstrax Tech Inc. we hereby declare that our Terpenes and Terpene Blend products are suitable for consumption by:

Vegans (do not eat meat, poultry, fish/seafood, eggs dairy products, honey)

Lacto-Vegetarians (do not eat meat, poultry, fish/seafood, eggs, but do eat dairy products)

Ovo-Vegetarians (do not eat meat, poultry, fish/seafood, dairy products, but do eat eggs)

Lacto-Ovo-Vegetarians (do not eat meat, poultry, fish/seafood, but do eat eggs and dairy products)

Donald Remmel

Donald Themen

Quality Control Inventory Analyst



April 23, 2021

ABSTRAX TECH PRODUCTS AS PHARMACEUTICAL EXCIPIENTS

Excipients include flavors that are used to improve the patient acceptability of a pharmaceutical product. The Pharmaceutical and Food Industries acknowledge a gap between the GMP's of the two Industry's and also that it generally doesn't make financial sense for a food ingredient manufacturer to try to bridge this gap for a relatively small number of pharmaceutical customers. Instead, as long as a food ingredient manufacturer can show evidence of having "a documented quality system, including procedures, test methods, results, validation, and change control notification"¹, then they can generally supply to the pharmaceutical industry. Therefore, we would like to assure you that Abstrax is certified to ISO 9001:2015² with a documented quality system for the design, manufacture, distribution and research of terpene flavor blends. Further, Abstrax products are manufactured with FEMA-GRAS chemicals and in strict adherence to internal specifications for use as flavors for the food industry. According to the International Pharmaceutical Excipients Council (IPEC), reference to the FEMA GRAS evaluations can be included in the Drug Master File (DMF) to support the safe use of a particular flavor in a pharmaceutical application³. Although we believe that products produced by Abstrax may qualify for use as pharmaceutical excipients, we manufacturer our terpene flavor blends for use in foods and will not guarantee their compliance with pharmaceutical standards.

Abstrax is dedicated to providing our customers with safe, high quality products. Thank you for your continued interest in our terpene flavor blends.

Sincerely.

Donald Remmel

QC/Inventory Analyst

1 USP FCC 7, General Information / Food Ingredients: Pharmaceutical Applications and Use of Appropriate GMPs

2 https://cdn.shopify.com/s/files/1/2149/6413/files/Abstrax Tech Inc. final cert.pdf?v=1597110558

3 Pharmaceutical Technology, June 2002, The IPEC-Americas Excipient Master File Guide, Christopher DeMerlis



15550 Rockfield Blvd., Suite B120, Irvine, California 92618

September 6, 2019

VITAMIN E ACETATE

Abstrax Tech, Inc. has not and does not use Vitamin E Acetate in the manufacture of any of its products.

Kevin Koby,

Chief Science Officer

PHYTOL STATEMENT



2661 Dow Avenue, Tustin, CA 92780

April 27, 2021

Phytol Statement For Terpene Blends

At Abstrax Tech Inc. we use very specific ingredients in order to achieve the desired flavor and effect. We do not add any of the acyclic diterpenoid called Phytol to any of our blends.

Donald Remmel

Sould Hum

QC/Inventory Analyst



Updated March 11, 2021

Abstrax Statement on Vape Cartridge Ingredients

Abstrax has been diligently following the CDC's investigation of E-cigarette, or Vaping. use-Associated Lung Injury (EVALI)1. The CDC has concluded that Vitamin E Acetate is the primary cause of EVALI as it was identified in >94% of the lung samples taken from studied EVALI patients. Toxicants from diluent terpenes were only found in 1 out of 51 samples from EVALI patients. Therefore, Abstrax would like to assure our customers that we have never used Vitamin E Acetate in any of our formulations. Furthermore, if we become aware of conclusive evidence that any ingredients we use will pose a risk to the consumer when used in vaporizers, e-cigarettes, and electronic nicotine delivery systems (END), then our customers will be informed immediately. Abstrax will continue monitoring scientific reports to assure the safety of our products. EVALI cases peaked in August 2019 and have persistently declined ever since. The CDC suspects that the decline could be related to removal of Vitamin E Acetate from products as well as increased public awareness and law enforcement actions against illicit END products purchased off the street or from unknown sources. EVALI has proven that it is essential for consumers to rely on reputable suppliers to satisfy their END indulgence. Inhalable products have been a rapidly growing segment but there is currently a paucity of research about the safety of these products. Reputable suppliers are more likely to consider all available evidence to try and assure the safe use of ingredients they market for applications leading to inhalation. An integral part of vaping is the composition of the vape liquid, and some type of diluent or carrier is necessary for a satisfactory vaping experience. Abstrax Tech offers TEC-Temper as our proprietary, flavored diluent/carrier ingredient. This is composed of naturally-sourced substances that perform well in a variety of applications. Based on available data, Abstrax believes that TEC-Temper can be safely used in topicals, foods, beverages also devices described as vaporizers, ecigarettes, and electronic nicotine delivery systems (END). Specifically concerning inhaled application use, we base our conclusion regarding safety, in part, on a history of safe use in other inhaled applications such as fragrances for perfumes, aerosol deodorants, candles, incense and heated oil plug-in air fresheners. All the ingredients used in TEC-Temper have been evaluated by the Research Institute for Fragrance Materials (RIFM) and/or the Cosmetic Ingredient Review (CIR) for safe use in fragrances_{2,3}. However this alone does not guarantee the safety of our products when inhaled. Abstrax has proactively challenged itself to address the lack of safety data for inhaled ingredients. Last year, Abstrax initiated a research collaboration with the University of California at Riverside (UCR) to study the safety and efficacy of substances used as carriers, diluents, and stabilizers in the formulation of inhalable products. We have included our own products in these evaluations. At present we do not have results that we can share as we are completing a preliminary study. We will follow this with additional studies to better understand the effects that these substances may have when inhaled.



Abstrax is a team of scientists who care deeply not only about our customers and our community, but also our industry and the safety and efficacy of our products. Once we have completed our studies and properly evaluated the results through the peer review process, we intend to share this information publicly. We want everyone to be equally informed so they can make educated decisions about the products they purchase and/or consume. In sharing this information, we are taking a position of safety over profits. We understand there is a chance that we may not like the results of our studies, however, we see it as critical for the safety of our customers and the community in general. Abstrax will continue to aggressively monitor and develop scientific data to assure the safety of our products. In the meantime, the FDA is recommending that consumers, "avoid buying vaping products of any kind on the street, and to refrain from using THC oil or modifying/adding any substances to products purchased in stores. If you continue to use these THC containing vaping products, monitor yourself for symptoms (e.g., cough, shortness of breath, chest pain) and promptly seek medical attention if you have concerns about your health."4

- 1. https://www.cdc.gov/tobacco/basic_information/e-cigarettes/severe-lung-disease.html
- 2. The safety of fragrances is assured by the Research Institute for Fragrance Materials (RIFM), a non-profit corporation that was formed in 1966 "to encourage"
- uniform safety standards related to the use of fragrance ingredients."; https://www.rifm.org/about.php
- 3. The Cosmetic Ingredient Review (CIR) was established in 1976, with the support of the U.S. Food and Drug Administration and the Consumer Federation of
- America, to review and assess the safety of ingredients used in cosmetics and publish the results in peer-reviewed scientific literature; https://www.cirsafety.
- 4. https://www.fda.gov/consumers/consumer-updates/vaping-illnesses-consumers-can-help-protect-themselves-avoiding-tetrahydrocannabinol-thc-containing

QC Inventory Analyst

Sound Hum

Whole Kosher Services

10819 Sandpiper Drive #182 Houston, TX 77096 (210) 913-1836



June 23, 2021 13 Tamuz, 5781

Certificate of Kashruth

To Whom it may Concern,

This is to certify that there was an inspection of the facility: <u>Abstrax Tech., Inc. Located at 2661 Dow</u> <u>Avenue, Tustin, CA 92780</u> We hereby certify that the following products are Kosher pareve (contain no dairy or meat) and may be used all year round excluding Passover.

This certificate is valid until June 23, 2022 at which time it is subject to renewal.

Signed _

Rabbi Yaakov Cohen Kashrut Administrator

Page 1 of 17

QUALITY STATEMENT

We are industry leaders in terpene science, not only from a research perspective, but also where product quality and safety are concerned. We are ISO 9001:2015 certified and in the process of being HACCP certified for total compliance and have the highest purity standards in the industry.

We adhere to rigorous processing SOPs and our products are 100% free of fillers, meaning our terpenes are undiluted by PG, VG, MCT, or coconut oil.

Abstrax conducts ongoing quality assurance procedures to maintain the highest standards and is consistently searching for ways to improve. With two Ph.D.'s on staff and 50 years of combined experience, we invest in extensive terpene research and development to bring you not only the highest quality terpenes with the most complex flavor profiles in an ever-growing selection to help you separate your products from your competition. To the best of our knowledge and ability to ascertain, our products meet all applicable federal, state, and local regulatory requirements for solvent residues, heavy metals, pesticides and other residues.

QUALITY POLICY

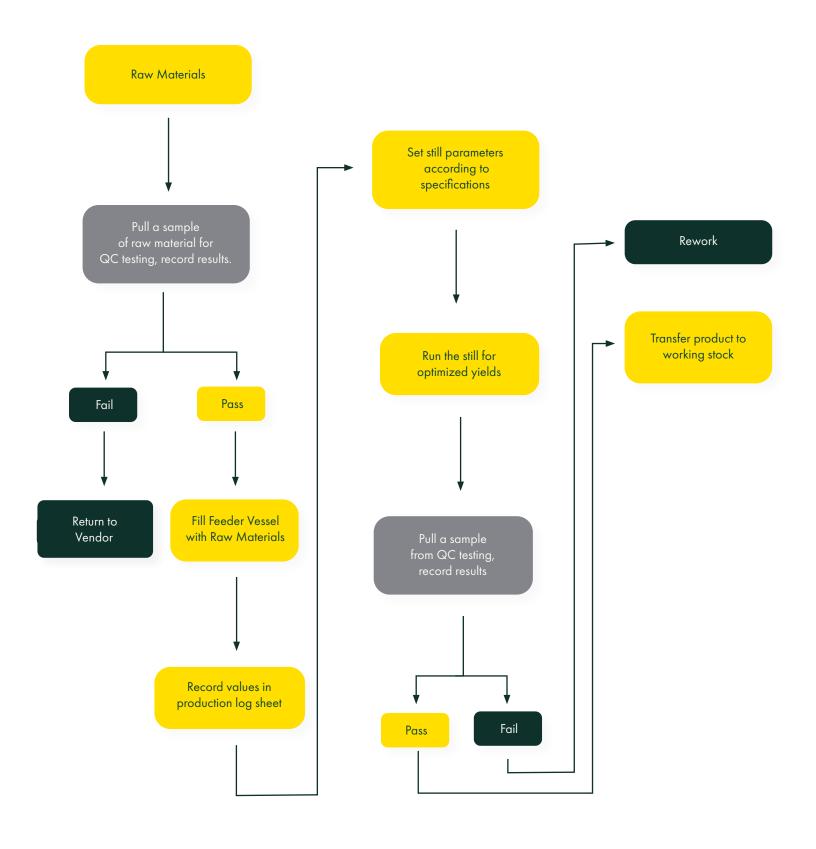
Abstrax Tech Inc. Made by Nature, Perfected by Science, provides high quality, naturally derived terpene blends. Product consistency and a high level of customer service are our core values. We achieve this by employing a highly skilled team dedicated to continuous improvement of our quality management system and ensuring that we meet our customer and statutory requirements.

Commitment to our customers

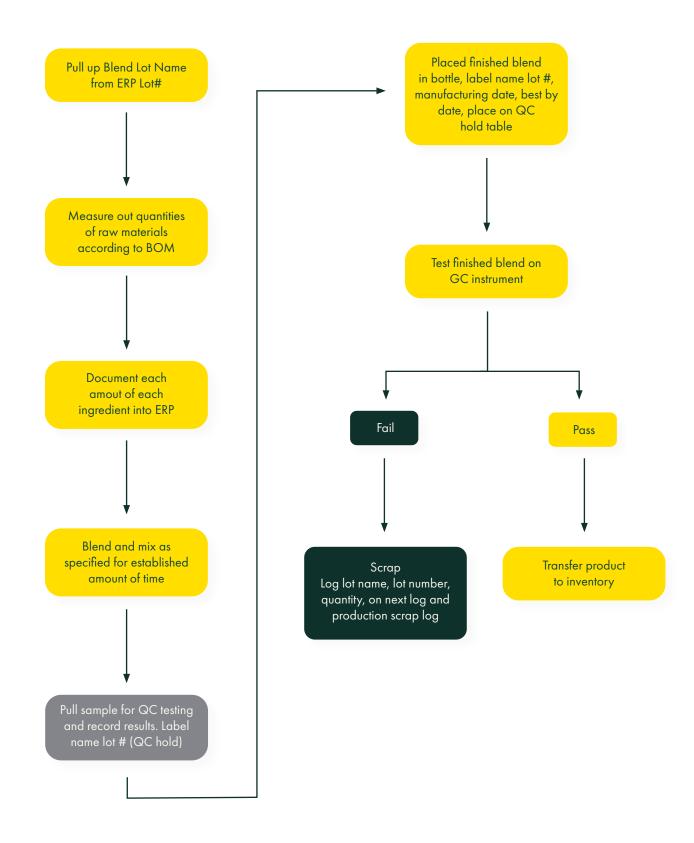
To ensure best practice, Abstrax Tech follows the internationally recognized Hazard Analysis Critical Control Point (HACCP) system and follows ISO 9001:2015.

To achieve our goal, we:

- Perform regular identification of hazards, determination of critical control points and timely implementation of effective control and monitoring measures
- Conform with the regulatory/statutory requirements and the agreed customer requirements
- Define the terpene safety objectives and continually review to ensure consistent compliance
- Communicate, implement and maintain this practice at all levels of the company
- Employ competent staff, reliable contractors and source from reputable suppliers
- Provide our personnel with adequate HACCP information, training, instructions, tools and equipment to carry out their job in a hygienic and professional manner
- Promote personal hygiene and cleanliness to our staff, contractors, suppliers and visitors
- Develop and strive to continually improve our processes capable of delivery of safe products through an efficient, effective and sustainable quality management system



30



General Personal Safety and Hygiene

- a. Employees are expected to show up for work in a clean, healthy and safe manner.
- b. Smoking is prohibited anywhere on the premises.
- c. No food is permitted in production areas, including but not limited to food, drinks, chewing gum/tobacco, candy, lozenges and cigarettes. Medication may be stored in personal lockers but is prohibited in production and warehouse areas.
- d. Personnel shall refrain from sneezing or coughing over materials or products.
- e. Clean uniforms and lab coats must be worn in production areas.
- f. Clean closed-toed shoes must be worn. Long hair is secured and within hair net. No jewelry is allowed.
- g. Personal lockers are provided and must be used for personal property only. Lockers are separated from production areas.
- h. The following is Performed in the Production Area only:
- i. Perform pre-operational check at the beginning of each workday. (Equipment check)
- j. Before entering the Manufacturing area, be sure to pay attention to the PPE signs and don PPE accordingly.
- k. Always wear gloves and eye protection when working with raw materials.
- I. All PPE is provided such as goggles, masks, hairnets, beard nets, and gloves.
- m. Clean working stations before and after task completion.
- n. If a spill occurs on the working stations, report the spill to management and clean up immediately.
- o. Hand Washing
 - 1. Proper hand washing steps are: Rinse hands; Apply soap; Scrub and lather soap for 19 seconds; Rinse hands thoroughly; and Dry with a paper towel. Following hand washing. Using hand sanitizer DOES NOT replace proper hand washing.
- p. Hands must be washed:
 - 1. At the start of each shift (at start-up, after lunch and breaks)
 - 2. After using the bathroom or smoking
 - 3. After blowing nose, coughing, sneezing, etc
 - 4. After picking up items from the floor
 - 5. Any time your hands become contaminated (touch dirty surfaces, garbage bins, etc.)
 - 6. When entering the production area from a less-clean area (e.g. outside or warehouse)
- q. Illness: If an employee has experienced symptoms of an infectious disease (i.e. diarrhea, vomiting, sores/wounds, sore throat, fever, etc.) within the last 24 hours, the employee shall report illness to management and shall be prohibited to work and sent home by his/her supervisor to protect the other employees and the safety of the food. Personnel with wounds or burns shall be required to cover them with brightly colored or metal detectable dressings if in the production area. Any lost dressing shall be reported to management immediately.

* A check mark (②)indicates the Allergen/Sensitive Agent is present. If blank, it means that to the best of our knowledge, there are no Allergen / Sensitive agents present.

Allergen/Sensitive Agent	Source of Allergen in the Product*	Present in Product*	Present in the Same Line*	Present in the Facility*
CORN & CORN PRODUCTS (Includes modified starch, hydrolyzed protein, sweeteners, sugars, spice carriers)	\bigcirc	\bigcirc	\bigcirc	\bigcirc
EGG & EGG PRODUCTS (liquids and powders)	\bigcirc	\bigcirc	\bigcirc	\bigcirc
FISH & FISH PRODUCTS (Includes any and all species of fresh and saltwater fish)	\bigcirc	\bigcirc	\bigcirc	\bigcirc
SEEDS	\bigcirc	\bigcirc	\bigcirc	\bigcirc
GLUTEN (Wheat, rye, barley, oats, flour, etc.)	\bigcirc	\bigcirc	\bigcirc	\bigcirc
MILK & DAIRY PRODUCTS (Includes whey, lactose, cheese, casein, spice carriers, milk, cream, etc.)	\bigcirc	\bigcirc	\bigcirc	\bigcirc
FD&C Yellow 5	\bigcirc	\bigcirc	\bigcirc	\bigcirc
PEANUTS, PEANUT OIL & PEANUT DERIVED ITEMS (Peanut meal, flour & ground nuts, szechuan sauce, mandelona nuts, etc.)	\Diamond	\bigcirc	\bigcirc	\bigcirc
SESAME SEEDS & SESAME OIL	\bigcirc	\bigcirc	\bigcirc	\bigcirc
CEREAL PROTEINS (Including HVP)	\bigcirc	\bigcirc	\bigcirc	\bigcirc
SHELLFISH, CRUSTACEANS & MOLLUSCS	\bigcirc	\bigcirc	\bigcirc	\bigcirc
SOY (except refined soy oil)	\bigcirc	\bigcirc	\bigcirc	\bigcirc
SULFITES (Includes sulfur dioxide, sodium dithionite, chemicals that lists sulfite, etc.)	\bigcirc	\bigcirc	\bigcirc	\bigcirc
TREE NUTS (Includes almonds, beechnuts, brazil nuts, nutmeg, cashews, chestnuts, coconut, etc.)	\bigcirc	\bigcirc	\bigcirc	\bigcirc
WHEAT, GLUTEN & TRITCALE	\bigcirc	\bigcirc	\bigcirc	\bigcirc
MUSTARD & MUSTARD OIL	\bigcirc	\bigcirc	\bigcirc	\bigcirc
AUTOLYZED YEAST	\bigcirc	\bigcirc	\bigcirc	\bigcirc
CELERY	\bigcirc	\bigcirc	\bigcirc	\bigcirc

There are currently no allergens on-site or in the products.

GAS CHROMATOGRAPHY ANALYSIS COMPOUND TEST

We perform Gas Chromatography analysis for two reasons: (1) to ensure that the terpene blends produced contain the correct percentages of their isolates & (2) to determine that any residual solvents used in the manufacture & refinement of our raw materials are either absent from our products or present in quantities below the legal thresholds. The state of California requires testing for the 22 following specific compounds:

Category I Residual Solvent	CAS No.	Cannabis Product Action Level PPM
1,2-Dichloromethane	107-06-2	1.0
Benzene	71-43-2	1.0
Chloroform	67-66-3	1.0
Ethylene oxide	75-21-8	1.0
Methylene chloride	75-09-2	1.0
Trichloroethylene	79-01-06	1.0
Category II Residual Solvent	CAS No.	Cannabis Product Action Level PPM
Acetone	67-64-1	5000
Acetonitrile	75-05-8	410
Butane	106-97-8	5000
Ethanol	64-17-5	5000
Ethyl acetate	141-78-6	5000
Ethyl ether	60-29-7	5000
Heptane	142-82-5	5000
Hexane	110-54-3	290
Isopropyl alcohol	67-63-0	5000
Methanol	67-56-1	3000
Pentane	109-66-0	5000
Propane	74-98-6	5000
Toluene	108-88-3	890
Total xylenes (ortho-meta-para)	13330-20-7	2170

If the sample fails residual solvents and processing chemicals testing, the batch from which the sample was collected fails residual solvents and processing chemicals testing and shall not be released for retail sale.

GAS CHROMATOGRAPHY ANALYSIS COMPOUND TEST (CONT.)

Comments

- 1. CA Residual Solvents Limits are applied to undiluted raw materials (isolates)
- 2. CA Residual Solvents Limits are applied to undiluted Finished Products (terpene flavor profiles.)

Document Prepared: <u>Donald Remmel</u>	
Signature: Novald Human	Date: <u>01-03-2020</u>
Document Verified: Winston A. Boyd, Ph.D.	
Signature:	_ Date: <u>01-03-2020</u>
Document Approved: <u>Donald Remmel</u> Signature:	_ Date: <u>01-03-2020</u>

Question 1: What certifications does Abstrax Tech possess?

We are currently ISO 9001:2015 and HACCP certified. We follow the strictest limits across 50 states when analyzing each raw material and each finished product lot for safety (Residual Solvents, Pesticides and Heavy Metals.)

Question 2: What is ISO 9001:2015?

ISO 9001:2015 (International Standard Organization) specifies requirements for a quality management system when an organization:

- a. Needs to demonstrate its ability to consistently provide products and services that meet customer and applicable statutory and regulatory requirements, and
- b. Aims to enhance customer satisfaction through the effective application of the system, including processes for improvement of the system and the assurance of conformity to customer and applicable statutory and regulatory requirements.

All the requirements of ISO 9001:2015 are generic and are intended to be applicable to any organization, regardless of its type or size, or the products and services it provides.

Question 3: What is cGMP?

Current Good Manufacturing Practice (GMP) is a system for ensuring that products are consistently produced and controlled according to quality standards. It is designed to minimize the risks involved in any pharmaceutical production that cannot be eliminated through testing the final product. cGMP is performed in accordance with HACCP.

Question 4: What is HACCP?

HACCP stands for Hazard Analysis and Critical Control Points. This is a preventative food safety system in which every step in the manufacture, storage and distribution of a food product is scientifically analyzed for microbiological, physical and chemical hazards.

Question 5: What is ISO/IEC 17025?

We use an ISO Certified Lab for testing our raw material CBD.

ISO/IEC 17025 General requirements for the competence of testing and calibration laboratories is the main ISO standard used by testing and calibration laboratories. In most countries, ISO/IEC 17025 is the standard for which most labs must hold accreditation in order to be deemed technically competent.

Question 6: Why does certification matter?

Certification shows that the company has adequately demonstrated to a third-party that it meets the requirements of a certain standard and is dedicated to continuous improvement, managing risk, and maintaining customer satisfaction. The result of an effective quality system is consistent, safe, and quality products.

FREQUENTLY ASKED QUESTIONS (CONT.)

Question 7: Do you have a Recall Plan?

Yes, it is a part of our HACCP Plan. Mock recalls are performed annually. We have total traceability from bulk materials to every product sent to every customer.

Question 8: What documents are available on the website?

COA (Certificate of Analysis), SDS (Safety Data Sheets), and PDS (Product Data Sheets).

Question 9: Do you have regulatory registrations, liability insurance, etc?

Yes, we have the following documents: current FDA registration, Liability Insurance. These documents are available per request.



www.abstraxtech.com | info@abstraxlabs.com | 2661 Dow Avenue | Tustin, CA 92780





Name: Natural Mimosa Matrix:Terpene Blend Lot #:1370

Product #: ABXTB0098

Manufacture Date: 4/27/2021 Expiration Date: 4/27/2022

Overall Result: Pass

Test	Specification	Result	
Appearance	Colorless to yellow clear liquid	Pass	

Volumetric glassware complies with Class A tolerance requirements of ASTM E 288 and NIST Circular 434; it is calibrated before first use and recalibrated regularly in accordance with ASTM E 542 and NIST Procedure NBSIR 74-461. Balances are calibrated regularly with weights certified traceable to the NIST national mass standard. Thermometers and temperature probes are calibrated before first use and recalibrated regularly with a thermometer traceable to NIST standards. All products are prepared according to master documents that assure manufacture according to validated methods. Batch records document raw material traceability and production and testing history for each lot manufactured.

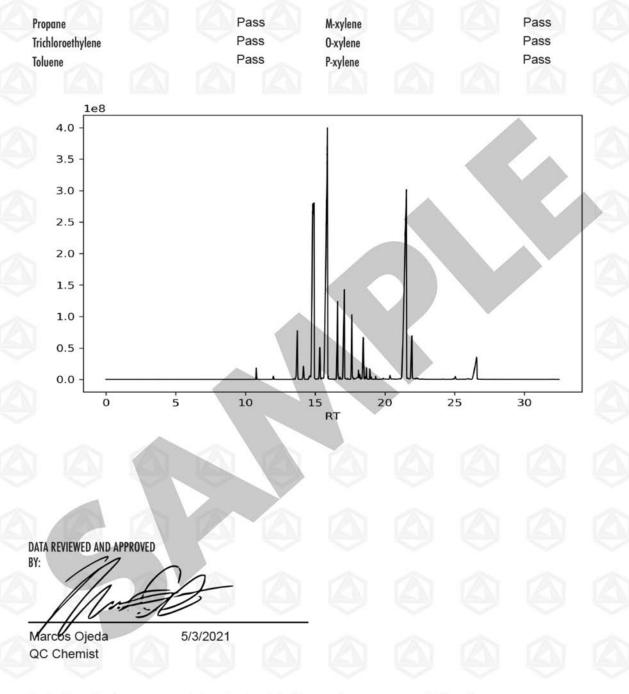
Specific Gravity @ 25 °C	Density Range	Result
0.8596	0.84-0.88	Pass

Gas Chromatograph	Residual Solvent Analysis	Water Activity (A _W) < 0.65	
>90% purity	Pass	0.4	

Pass	Ethyl Ether	Pass
Pass	Ethylene Oxide	Pass
Pass	Heptane	Pass
Pass	Hexane	Pass
Pass	Isopropyl Alcohol	Pass
Pass	Methanol	Pass
Pass	Methylene Chloride	Pass
Pass	Pentane	Pass
	Pass Pass Pass Pass Pass Pass	Pass Ethylene Oxide Pass Heptane Pass Hexane Pass Isopropyl Alcohol Pass Methanol Pass Methylene Chloride

www.abstraxtech.com | info@abstraxlabs.com | 2661 Dow Avenue | Tustin, CA 92780

CERTIFICATE OF ANALYSIS (CONT.)



This Certificate of Analysis is accurate to the best of our knowledge. However, the customer is responsible for performing its own assessment to ensure that the material is suitable for its intended applications, including compliance with all pertinent legal requirements. The expected shelf life is calculated from the original manufacture date and is based on unopened containers stored under proper conditions. Once containers have been opened, maintaining minimal headspace and storing under appropriate conditions will aid in preserving the product's integrity.

www.abstraxtech.com | info@abstraxlabs.com | 2661 Dow Avenue | Tustin, CA 92780



www.abstraxtech.com | info@abstraxtech.com | 2661 Dow Avenue, Tustin, CA 92780

SAFETY DATA SHEET

Originally Prepared: 06/21/2019

Version: 1.0

1. IDENTIFICATION

1.1 Product Identification:

Concentrated Flavor Mixtures including Natural Animal Mints Rosin (Native Series) PRODUCT NUMBER:

ABXTB0330H, ABXTB0330G, ABXTB0330E, ABXTB0330C, ABXTB0330A, ABXTB0330D, ABXTB0330F, ABXTB0330Y, ABXTB0330X

CAS NUMBER: Not applicable. EINECS NUMBER: Not applicable. E NUMBER: Not applicable.

1.2 Product Use

Concentrated flavor mixture for foodstuffs in which added flavoring is permitted.

1.3 Company Identification

Abstrax Tech Inc. 15550 Rockfield Blvd. Suite B, Irvine, CA. 92618, USA; 1-562-294-5805

www.abstraxtech.com

SDS inquiries can also be submitted to regulatory@abstraxtech.com

1.4 Emergency Telephone Number

USA 1-562-294-5805 (only during office hours, 8:00am - 5:00pm PST)

2. HAZARD(S) IDENTIFICATION

2.1 Classification of the Product

This mixture is not classified as dangerous. The flavor poses no hazards under normal conditions of handling and usage. If misused in its concentrated form, the flavor may be harmful to health.

2.2 GHS Label Elements

Pictogram:



Signal Word: Warning

Hazard Statement(s): H227 Flammable liquid and vapor

Precautionary Statement(s):

Prevention:

P262 Do not get in eyes, on skin, or on clothing.

P264 Wash hands thoroughly after handling.

P272 Contaminated work clothing must not be allowed out of the workplace.

P280 Wear protective gloves, clothing and eye protection.

P302+P352 - IF ON SKIN (or hair): Wash with plenty of water.

P304 + P341 IF INHALED: If breathing is difficult, remove to fresh air and keep at rest in a position comfortable for breathing.

P305 + P338 +P351 IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing.

P309 + P311 IF you feel unwell, call a poison center or physician.

P333 + P313 If skin irritation or rash occurs: Get medical advice/attention.

P342 + P311 If experiencing respiratory symptoms: Call a POISON CENTER or physician.

SDS Page 1 of 5

P337 + P313 If eye irritation persists: Get medical advice/attention.

P361 Remove immediately all contaminated clothing.

P363 Wash contaminated clothing before reuse.

P370 In case of fire: Use CO2, Foam, Dry Chemicals, Water for extinction.

Storage

P403 + P235 Store in a well-ventilated place. Keep cool.

Disposal:

P501 Dispose of contents and container in accordance with local, regional, national and international regulations.

2.3 Other Hazards

Exposure may aggravate those with pre-existing eye, skin, or respiratory conditions.

2.4 Acute Toxicity (GHS-US)

No data available. This flavor mixture has unknown acute toxicity. However, this mixture consists entirely of ingredient(s) which are approved for use in a regulation of the Food and Drug Administration (FDA) and/or are regarded as safe by a reliable industry association.

3. COMPOSITION / INFORMATION ON INGREDIENTS

3.1 This product is a concentrated flavor mixture. All flavor ingredients contained in this product are approved for use in a regulation of the Food and Drug Administration (FDA) or are regarded as safe by a reliable industry association. Specific chemical identities and/or exact percentages of composition are being withheld as a trade secret pursuant to 29CFR §1910.1200.

CAS NUMBER: Not applicable. EINECS NUMBER: Not applicable.

4. FIRST-AID MEASURES

4.1 General First-Aid Instructions

INHALATION: If headache, irritation, nausea, or drowsiness occurs, move patient to a place with clear air. Ventilate. Obtain medical advice if irritation persists.

SKIN CONTACT: Remove contaminated clothing. Wash with water and soap. Obtain medical advice if irritation persists. An eyewash and safety shower should be readily available.

EYE CONTACT: Check for and remove any contact lenses. Flush opened eye (including under eyelids) with running water for 15 minutes. Hold eyelids apart while flushing to rinse entire surface of eyes and lids. Do not allow person to rub eyes or keep eyes closed. Obtain medical advice if irritation persists. An eyewash and safety shower should be readily available.

INGESTION: If patient is conscious and can swallow, administer lots of water to dilute substance. Do not induce vomiting. Call a poison control center and obtain medical advice if irritation persists.

OTHER: Personal protective equipment (PPE) for first aid responders is recommended; Chemical resistant clothing, boots and gloves.

4.2 Most Important Symptoms and Effects, Acute and Delayed

Concentrated flavor may initially irritate eyes, lungs, skin and stomach. Call a poison control center and obtain medical advice if irritation persists after following above instructions.

4.1 Immediate Medical Care and Special Treatment

Call a poison control center and obtain medical advice if irritation persists.

5. FIRE-FIGHTING MEASURES

5.1 SUITABLE EXTINGUISHING MEDIA: CO2, Foam, Dry Chemicals, Water Spray

NON-SUITABLE EXTINGUISHING MEDIA: Powder, hazards of dust cloud formation. Do not use a heavy water stream. A heavy water stream may spread burning liquid.

5.2 SPECIFIC FIRE AND EXPLOSION HAZARDS: Heat may generate irritating or corrosive vapors. Product is a flammable liquid and considered to be a fire or explosion hazard.

HAZARDOUS COMBUSTION PRODUCTS: Carbon dioxide, carbon monoxide, fumes.

5.3 SPECIAL PROTECTIVE EQUIPMENT OR PRECAUTIONS FOR FIREFIGHTERS: Exercise caution when fighting any chemical fire. Use protective clothing and breathing equipment appropriate for surrounding fire. Prevent contact with skin and eyes.

6. ACCIDENTAL RELEASE MEASURES

PERSONAL PRECAUTIONS: Keep concentrate away from heat, sparks, open flames, hot surfaces. No smoking. Ventilate site. Evacuate unnecessary personnel. Wear safety glasses or goggles, rubber chemical resistant gloves, and apron when handling concentrate. Pressure demand air supplied respirators should always be worn when the airborne concentration of the contaminant or oxygen is unknown. Otherwise, wear NIOSH approved respiratory protection and other personal protective equipment as appropriate for the potential exposure hazard. Do not get into eyes, on skin, or clothing.

METHODS AND MATERIALS FOR CONTAINMENT: Contain large spills with dikes or absorbents to prevent migration and entry into sewers or streams.

METHODS AND MATERIALS FOR CLEAN UP: Absorb large spills with vermiculite or other suitable absorbent material. Shovel up material and place in approved disposal container. Large spills may also be pumped into closed containers for recovery and disposal. Dispose of in

SDS Page 2 of 5

accordance with current local, state, and federal laws and regulations. Flush the area with warm water. Small spills may be flushed to the drain with excess water or absorbed with a damp cloth.

7. HANDLING AND STORAGE

7.1 Handling

Maintain good ventilation. Consistent with good hygiene and lab practices, wear safety glasses or goggles, rubber gloves and apron when handling. Where aerosols of concentrate are created, use suitable dust mask or breathing apparatus. Do not eat, drink or smoke in work areas. Remove contaminated clothing and protective equipment before entering eating areas. Avoid prolonged or repeated exposure to concentrate. Wash thoroughly after handling.

7.2 Storage

Keep container tightly sealed. Store in cool, dry, secure, flammable storage area away from all sources of ignition. Protect from light, avoid uncoated metal containers, and keep air contact to a minimum. Incompatible with strong oxidizing agents, acids and bases. Avoid extended storage by using as soon as possible within designated shelf life.

7.3 Specific Uses

Concentrated flavor mixture for foodstuffs in which added flavoring is permitted. It is the user's responsibility to ensure that the conditions and possible uses of the flavor conform to local laws and regulations.

8. EXPOSURE CONTROLS / PERSONAL PROTECTION

8.1 Exposure Limit Values

There are no established exposure limits for this mixture. This flavor poses no exposure hazards under normal conditions of handling and approved levels of ingestion. If misused in its concentrated form, the flavor may be irritating to the skin and eyes. Vapors or mist, in unusually high concentration generated from spraying, heating the material or as from exposure in poorly ventilated areas or confined spaces, may cause irritation of the throat and nose, headache, nausea and drowsiness.

Control Parameters: Not Applicable.

8.2 Exposure Controls

VENTILATION AND OTHER ENGINEERING CONTROLS: Normal ventilation is generally adequate. Keep airborne concentration of flavor low by enclosing production processes whenever possible.







EYE: Wear protective glasses, goggles or face shield. An eyewash and safety shower should be readily available.

HAND: Wear protective chemical resistant gloves during manufacture or handling of this concentrated flavor.

SPECIAL CLOTHING: Special clothing is not normally necessary when manufacturing or handling this flavor within a closed system. Wear chemical resistant full body coveralls whenever there is a possibility of splash or appreciable contact with vapors during manufacturing or handling of the concentrate.

OTHER PROTECTIVE DEVICES AND PROCEDURES: Follow good manufacturing practice.

RESPIRATORY: Work in well ventilated area. If irritation is experienced, approved respiratory protection should be worn.

9. PHYSICAL AND CHEMICAL PROPERTIES

9.1 General Information

APPEARANCE: Colorless to yellow, clear liquid ODOR: Pine, Citrus, Mint, Earthy ODOR THRESHOLD: Data not available

9.2 Important Health, Safety and Environmental Information

pH: Not applicable
MELTING POINT: Not applicable
FREEZING POINT: Data not available
BOILING POINT: Data not available
FLASH POINT: >73°F (23°C); <140°F (60°C)

FLAMMABILITY: Not applicable

EXPLOSIVE PROPERTIES: Data not available OXIDISING PROPERTIES: Data not available VAPOR PRESSURE: Data not available

RELATIVE DENSITY (SPECIFIC GRAVITY @ 25°C):

0.84-0.88 SOLUBILITY: Oil

WATER SOLUBILITY: Insoluble

PARTITION COEFFICIENT: Data not available

VISCOSITY: Data not available VAPOR DENSITY: Data not available EVAPORATION RATE: Data not available

9.3 Other Information

MISCIBILITY: Data not available

FAT SOLUBILITY (VEGETABLE OIL): Soluble

CONDUCTIVITY: Data not available

AUTO-IGNITION TEMPERATURE: Data not available DECOMPOSITION TEMPERATURE: Data not available

VOC CONTENT: Data not available

SDS Page 3 of 5

10. STABILITY AND REACTIVITY

10.1 Reactivity

Stable and non-reactive under normal conditions of use and storage. May form flammable/explosive vapour-air mixture in concentration.

10.2 Chemical Stability

This product is stable under normal storage conditions. Hazardous polymerization is not expected to occur. Avoid extended storage by using as soon as possible within designated shelf life.

10.4 Conditions to Avoid

Keep cool and avoid excessive heat, open flames, sparks or other sources of ignition. Avoid opening in poorly ventilated spaces.

10.5 Materials to Avoid

Avoid strong acids, bases, and oxidizing agents.

10.6 Hazardous Decomposition Products

No known hazardous materials produced in dangerous amounts upon decomposition. Thermal decomposition generates potentially irritating vapors.

11. TOXICOLOGICAL INFORMATION

This flavor mixture poses no known hazards under recommended conditions of handling and usage. If misused in its concentrated form, the flavor may be harmful to health. All flavor ingredients contained in this product are approved for use in a regulation of the Food and Drug Administration (FDA) or are regarded as safe by a reliable industry association.

Acute Toxicity (Oral, Dermal, Inhalation Exposure): Unknown.

Effects of Immediate, Delayed or Chronic Effects from Short- and Long-Term Exposure: Concentrate may cause immediate irritation to eyes, stomach, skin and/or lungs. Prolonged exposure may increase irritation. Data is not available regarding long-term effects of repeated exposures.

International Agency for Research on Cancer (IARC) Monograph: Mixture not evaluated.

OSHA Carcinogen: Mixture not evaluated. NIOSH Carcinogen: Mixture not evaluated.

National Toxicology Program (NTP) Report on Carcinogens (ROC): Mixture not evaluated.

12. ECOLOGICAL INFORMATION

12.1 Ecotoxicity

Not established.

12.2 Persistence and Degradability

Not established.

12.3 Bioaccumulative Potential

Not established. 12.4 Mobility in Soil Not established.

12.5 Other Adverse Effects

Not established. Avoid release into the environment.

13. DISPOSAL CONSIDERATIONS

Dispose of this flavor in accordance with Federal, State, County, and local regulations. Avoid any spillage from entering waterways, drains or sewage as this product has unknown ecotoxicity.

Containers should be either reconditioned by certified firms or properly disposed of by approved firms. Disposal of containers should be in accordance with applicable laws and regulations. "Empty" drums should not be given to individuals. Misuse of empty containers can be hazardous. Empty containers can be hazardous if used to store toxic, flammable, or reactive material. Cutting or welding of empty containers may cause fire, explosion, or toxic fumes from residues. Do not pressurize or expose to open flame or heat. Keep containers closed and drum bungs in place.

14. TRANSPORT INFORMATION

This flavor is classified as a Flammable Liquid based on the FP criteria of the GHS and US Department Of Transportation (DOT) regulations.

MODE: Ground, air, or vessel

UN SHIPPING NAME: Extracts, flavoring, liquid UN IDENTIFICATION NUMBER: UN1197

DOT HAZARD CLASS: 3 PACKING GROUP: III GHS CATEGORY: 3 SIGNAL WORD: Warning

HAZARD STATEMENT: Flammable liquid and vapor

HAZARD PLACARD REQUIRED:

SDS Page 4 of 5



STORAGE SEGREGATION: Usual precautions for flammable liquids

15. REGULATORY INFORMATION

UNITED STATES OF AMERICA (USA): All flavor ingredients contained in this product are approved for use in a regulation of the Food and Drug Administration (FDA) or are regarded as safe by a reliable industry association.

CALIFORNIA PROPOSITION 65:

WARNING: This product can expose you to Myrcene, which is known to the State of California to cause cancer. For more information go to www.P65Warnings.ca.gov.

16. OTHER INFORMATION

- 16.1 VERSIONS OF THIS DOCUMENT DATED PREVIOUS TO THE REVISION DATE ON THIS DOCUMENT ARE NO LONGER VALID.
- 16.3 THE ABOVE INFORMATION WAS PREPARED FROM CURRENT AND REPUTABLE SOURCES. HOWEVER, THE DATA IS PROVIDED WITHOUT ANY WARRANTY, EXPRESSED OR IMPLIED, REGARDING ITS CORRECTNESS OR ACCURACY.
- 16.5 TO AVOID RISKS TO HUMAN HEALTH AND THE ENVIRONMENT, COMPLY WITH THE INSTRUCTIONS FOR USE.
- 16.6 THE INFORMATION CONTAINED IN THIS DATA SHEET DOES NOT CONSTITUTE AN ASSESSMENT OF WORKPLACE RISKS. IT IS THE USERS' RESPONSIBILITY TO ENSURE SAFE CONDITIONS FOR HANDLING, STORAGE AND DISPOSAL OF THIS FLAVOR AND TO ASSUME LIABILITY FOR LOSS, INJURY, DAMAGE, OR EXPENSE RESULTING FROM IMPROPER USE OF THIS PRODUCT.

SAFETY DATA SHEET PREPARED BY ABSTRAX, JUNE 2019 IN CONFORMITY WITH OSHA 1910.1200



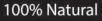


www.abstraxtech.com | info@abstraxtech.com | 2661 Dow Ave, Tustin, CA 92780

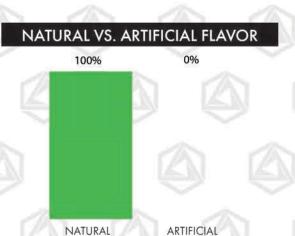
TERPENE SPEC SHEET Animal Mints Rosin (Native Series)

ANALYSIS DOMINANT TERPENES (>50%) TOP INGREDIENTS **D-Limonene** 20-50% **Beta Pinene** 5-10% Alpha Pinene 5-10% Linalool 5-10% Myrcene 1-5% MINOR TERPENES (<50%) Caryophyllene Camphene Ocimene Fenchol Alpha Caryophyllene Terpinolene **D-Limonene** Beta Pinene Alpha Terpineol **Bisabolene** Alpha Pinene Linalool Cadinene Alpha Terpinene Gamma-Terpinene p-Cymene Myrcene

Alpha Phellandrene Farnesene Bergamotene Borneol Guaiol Alpha Bisabolol Delta 3-Carene Sabinene Nerolidol 4-Terpineol Caryophyllene oxide Guaiene Citronellol L-Menthol Fenchone Eucalyptol



0% Artificial flavorings



"This information is proprietary commercial information and should be treated as a trade secret and kept confidential at all times



www.abstraxtech.com | info@abstraxtech.com | 2661 Dow Avenue, Tustin, CA 92780

Product Data Sheet

Animal Mints Rosin (Native Series)

Description:

Abstrax Animal Mints Rosinis an uplifting, creative blend with earthy, pine, citrus, mint, and skunkynotes.



Dominant Terpenes:

D-Limonene (20-50%), Beta Pinene (5-10%), Mycrene (10-20%), Alpha Pinene (5-10%), Linalool (5-10%), Myrcene (1-5%) and a blend of Alpha Caryophyllene, Sabinene and other minor terpenoids/flavor isolates

Mood Orientation: Uplifting, Creative

Specification (as manufactured):

Appearance	Colorless to yellow clear liquid		
Aroma	Pine, Citrus, Mint, Earthy		
Flavor	Pine, Citrus, Mint, Earthy		
Density	0.84-0.88		

www.abstraxtech.com | info@abstraxlabs.com | 2661 Dow Avenue, Tustin, CA 92780

Storage:

Store flavor in airtight, sealed containers at ambient temperature (<22°C/72°F) and <60% relative humidity. Avoid exposure to light, air and high heat. Once containers have been opened, maintaining minimal headspace and storing under appropriate conditions will aid in preserving the product's integrity.

Shelf Life:

Typical shelf life is 1 year from date of manufacture when stored in original container and under recommended storage conditions.

Food Ingredient Declaration For Customers

Natural Flovor (FDA)

This flavor contains ingredient(s) which are approved for use in a regulation of the Food and Drug Administration (FDA) and/or are listed as Generally Recognized as Safe (GRAS) on a reliable, published industry association list pursuant to 21CFR101.22.

Nutritional Contribution:

The nutritional content of flavor additives does not contribute to the nutritional value of foods, due to the extremely low use level. Therefore, pursuant to 21 CFR 101.9 (j) (4), flavors are exempt from nutritional labeling. Nutritional data is available upon request.

Regulatory Status:

Flavor			_		
USA Regulated Allergens					
International Regulated	Allergens ³	***			
Country of Maufacture		USA			
Other			Coi	ntains Myrcene	9

- 1. USA labeling recommendations are pursuant to 21CFR 101.22 (k) (2). Label declaration requirements vary significantly in different countries. These label declarations should not be used or adopted without confirming that the declaration is appropriate and legal for the subject country and intended application.
- 2. Crustacean, Egg, Fish, Milk, Peanut, Shellfish, Soybean, Tree Nut, Wheat are allergens recognized by the USA Food and Drug Administration Food Allergen Labeling and Consumer Protection Act of 2004 (FALCPA). FALCPA does not regulate sulfites as allergens. However, sulfites are considered a sensitizing chemical in the USA and will; be identified when present at >10 ppm.
- Apple, Beef, Buckwheat, Celery, Crustacean, Egg, Fish, Gelatine, Gluten, Kiwifruit, Milk, Mushroom, Mustard, Orange, Peanut, Pork, Poultry, Sesame Seed, Shellfish, Soybean, Sulfite, Tartrazine, Tomato, Tree Nut, Wheat, Yam
- WARNING: This product can expose you to Myrcene, which is known to the State of California to cause cancer. For more information go to www.P65Warnings.ca.gov.

www.abstraxtech.com | info@abstraxlabs.com | 2661 Dow Avenue, Tustin, CA 92780



For special inquiries, please contact:

Donald Remmel QC/Inventory Analyst qc@abstraxtech.com

Customer Service: info@abstraxtech.com