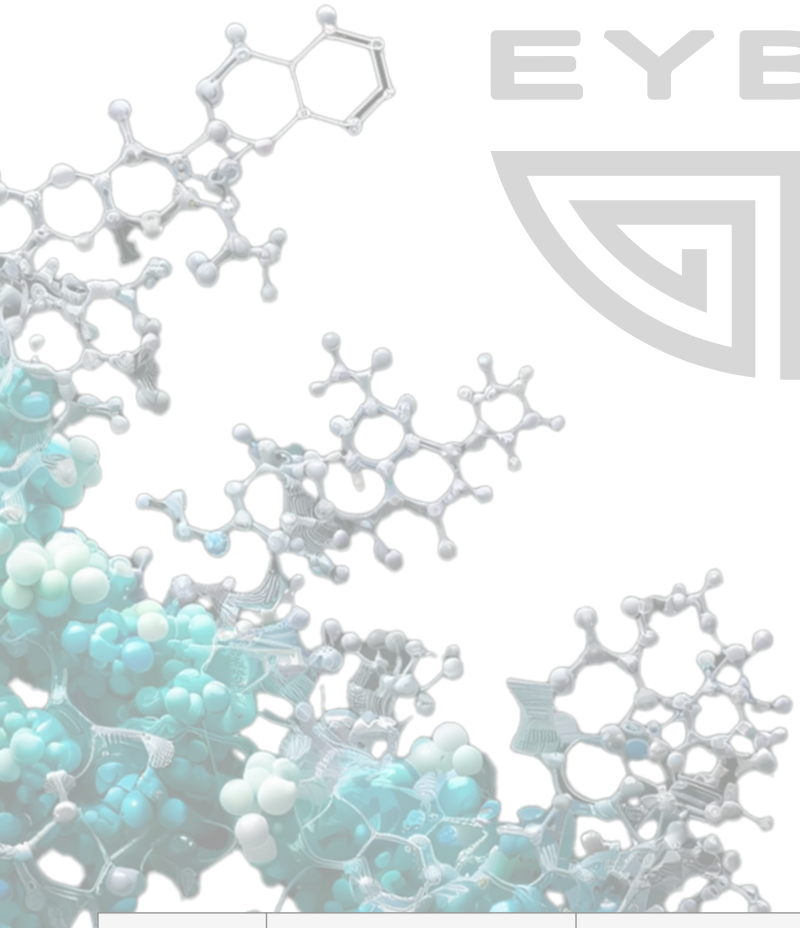
Compliance Packet

Eybna Technologies Ltd.



Version 03	**Last Updated:** June 6th, 2025 **Visit us:** www.eybna.com **Email:** info@eybna.com	**US Office** 7647 Hayvenhurst Ave #30. Van Nuys, CA 91406 USA +1 888 724 4813	**Israel Office** 1 Ha-Yozma St. Kfar Saba 4442214, Israel +972 3 3741976	**Berlin Office** Rheinsberger Str. 76/77. Berlin, 10115, Germany	

Table of Contents

1. Self Audit Form

General Information

Company Name	Eybna Technologies Ltd.
NIT Number	515146454
Mailing Address	1 Ha-Yozma St, Kfar Saba, 4442214, Israel
Phone Number	+972 3 3741976
Website	www.eybna.com
Email	info@eybna.com
Product Description or Service	Terpene-based formulations
Yaer of Establishment	2014
Number of Employees	40~
Number of Employees in Production	6
Number of Shifts	1
Square footage of the Facility	4305.56
QA Contact Name	Gil Tsapovetsky
QA Contact Email	gil.ts@eybna.com
Is Your Organization ISO 9001: 2015 Certified?	Yes
Is Your Organization FSSC 22000 Certified?	Yes
Is Your Organization GMP certified?	Yes

1. Quality Control System

1.1	Is there an established, maintained and documented quality system?	■ Yes □ No □ N/A
1.2	Is there a quality manual?	■ Yes □ No □ N/A
1.3	Is quality assurance independent of production?	■ Yes □ No □ N/A
1.4	Is the quality manual kept current and available to employees, auditors, or customers?	■ Yes □ No □ N/A
1.5	Are records kept of all products that are produced?	■ Yes □ No □ N/A
1.6	Do you have an internal audit program?	■ Yes □ No □ N/A
1.7	Are there formal written procedures for all performed product tests?	■ Yes □ No □ N/A
1.8	Do you use any contract laboratories?	■ Yes □ No □ N/A
1.9	Have you qualified/evaluated these contract laboratories?	■ Yes □ No □ N/A
1.10	What types of testing are contracted out?	Heavy Metals, Residual Solvents, Microbiological, Toxins, Pesticides
1.11	Are quality standards or written control procedures available for:	
1.12	Starting materials?	■ Yes □ No □ N/A
•	Finished products?	■ Yes □ No □ N/A
•	Are records kept of all control results?	■ Yes □ No □ N/A
1.13	If yes, for how long do you keep those records?	1 Year After Expiry
1.14	Is there a maintenance plan?	■ Yes □ No □ N/A
1.15	Do you analyse each sample?	■ Yes □ No □ N/A
1.16	Do you retain samples of each lot?	■ Yes □ No □ N/A
1.17	Do you have procedures covering the release or rejection of material?	■ Yes □ No □ N/A
1.18	Are batch records reviewed / approved before the batch is dispatched?	■ Yes □ No □ N/A
1.19	Are deviations and non-conformances investigated, documented and filed?	■ Yes □ No □ N/A
1.20	Would you notify your Customer of any deviations that occur during manufacturing?	Yes, when relevant

1. Quality Control System

1.21	Do you introduce changes according to a written procedure?	■ Yes □ No □ N/A
1.22	Do you inform your customers about changes?	Yes, when required
1.23	Are there procedures for the maintenance and use of laboratory equipment?	■ Yes □ No □ N/A
1.24	Do you have a documented glass and brittle plastic policy?	■ Yes □ No □ N/A
1.25	Are there documented standards for personnel hygiene?	■ Yes □ No □ N/A
1.26	Are there written and documented controls for contractors, visitors, agencies, or temporary staff?	■ Yes □ No □ N/A
1.27	Do you have an allergen cross-contamination control management program/policy?	■ Yes □ No □ N/A
1.28	Are your personnel instructed on handling any hazardous materials you use and how to act in case of unwanted events?	■ Yes □ No □ N/A

2. Personnel, Training and Education

2.1	Do you have written job descriptions for all personnel?	■ Yes □ No □ N/A
2.2	Do you have procedures that document how to perform training?	■ Yes □ No □ N/A
2.3	Do you maintain records of the training?	■ Yes □ No □ N/A
2.4	Does the training program in place have the following elements:	
2.5	Formal Introduction to Regulatory Guidance (ISO, etc.)	■ Yes □ No □ N/A
2.6	New Hire Program	■ Yes □ No □ N/A
2.7	Periodic assessment of practical effectiveness	■ Yes □ No □ N/A
2.8	Periodic refresher training programs for established employees	■ Yes □ No □ N/A
2.9	Training on new product manufacturing	■ Yes □ No □ N/A
2.10	Training on new methods that are used	■ Yes □ No □ N/A
2.11	Quality techniques for production people?	■ Yes □ No □ N/A

3. Facility and Utilities

3.1	Are there separate areas for:	
3.2	Handling of starting materials?	■ Yes □ No □ N/A
3.3	Manufacturing?	■ Yes □ No □ N/A
3.4	Quarantined finished products or are other control systems in place?	■ Yes □ No □ N/A
3.5	Approved finished products?	■ Yes □ No □ N/A
3.6	Packaging and dispatch?	■ Yes □ No □ N/A
3.7	Rest and eating?	■ Yes □ No □ N/A
3.8	Does the present design prevent:	
3.9	Chemical contamination?	■ Yes □ No □ N/A
3.10	Physical contamination?	■ Yes □ No □ N/A
3.11	Microbial contamination?	■ Yes □ No □ N/A
3.12	Are the working-rooms:	
3.13	Of proper size for the intended functions?	■ Yes □ No □ N/A
3.14	Satisfactorily lighted, air-conditioned?	■ Yes □ No □ N/A
3.15	Is there an adequate cleaning program in place?	■ Yes □ No □ N/A
3.16	Does the cleaning program employ analytical verification of cleanliness?	■ Yes □ No □ N/A
3.17	Supplied with security and fire protection measurements?	■ Yes □ No □ N/A
3.18	Are there written good housekeeping procedures?	■ Yes □ No □ N/A

4. Manufacturing

4.1	Are any penicillin, cephalosporin steroids, hormones, potent compounds, cytotoxins, insecticides, pesticides or other objectionable products manufactured at the facility?	□ Yes ■ No □ N/A
4.2	Is the material produced in a process that uses products of animal origin?	□ Yes ■ No □ N/A

4. Manufacturing

4.3	Is access to production area restricted?	■ Yes □ No □ N/A
4.4	Is adequate ventilation, air filtration and exhaust systems in place?	■ Yes □ No □ N/A
4.5	Is the control of re-circulated air sufficient to avoid contamination?	■ Yes □ No □ N/A
4.6	Is a preventive maintenance program in place?	■ Yes □ No □ N/A
4.7	Are there any raw materials in use that are derived from cannabis?	□ Yes ■ No □ N/A

5. Packaging, Labelling and Shipping

5.1	Is each bottle/container labelled with the lot/batch no.?	■ Yes □ No □ N/A
5.2	Will each bottle/container have the lot/batch no. and/or description clearly visible on it?	■ Yes □ No □ N/A
5.3	Do you keep records of all shipments to customers, including batch number and quantity?	■ Yes □ No □ N/A
5.4	Which cotractor are you using for transport shipping to customers?	UPS, FedEx, and DHL
5.5	If you use a contractor, do you have an agreed contract between parties which specifies required shipping conditions for materials?	■ Yes □ No □ N/A
5.6	If yes, have they been evaluated?	■ Yes □ No □ N/A
5.7	Are written instructions available for:	
•	Packaging components?	■ Yes □ No □ N/A
•	Packaging operation?	■ Yes □ No □ N/A
•	Labels and labelling?	■ Yes □ No □ N/A
5.8	Does the labelling procedure emphasize special precautions to prevent unintentional mix-up or substitution?	■ Yes □ No □ N/A
5.9	Do you maintain a lot separation during packaging?	■ Yes □ No □ N/A
5.10	Are you prepared to meet packaging and labelling requirements from your customers?	■ Yes □ No □ N/A

5. Packaging, Labelling and Shipping

5.11	Does your labelling indicate:	
•	Name and quantity?	■ Yes □ No □ N/A
•	The site of manufacturing?	■ Yes □ No □ N/A
•	The lot number?	■ Yes □ No □ N/A
•	Product code number?	■ Yes □ No □ N/A

6. Pest Control

6.1	Is pest control carried out by a third-party contractor?	■ Yes □ No □ N/A
6.2	Is the service contracted defined?	■ Yes □ No □ N/A
6.3	Is pest control carried out by trained personnel?	■ Yes □ No □ N/A
6.4	Are windows and doors to production areas adequately screened to prevent ingress of pets?	■ Yes □ No □ N/A
6.5	Are goods stored in such a way as to allow inspection and minimize the risk of infestation?	■ Yes □ No □ N/A
6.6	Are raw materials and finished products stored in clean, dry, and well-ventilated spaces, protected from dust, cross-contact, and sources of contamination?	■ Yes □ No □ N/A

2. Eybna's Plant Process Definitions

Raw materials

• Qualification is based on written specifications and approval of vendor sources.
• Reject/accept limits are defined by QA team.
• Approval is based on: Quality history, own QC, certificate and/or testing.
• Purity standard of raw material is 95% purity and above.
• Raw materials are visibly marked as approved.
• Raw materials are extracted by steam distilled and further purified by fractional distillation.
• ERP managed via Priority-ERP system and updated on daily basis.
• The storage area is separated and segregated.
• A stock rotation FEFO is in use.
• Rejected materials are clearly identified and physically segregated.
• Distinct staging area exists for raw materials.
• Research has shown that many of the molecules we use are also naturally found in cannabis. In specific product lines, like the Pure Terpenes line, every ingredient is confirmed to be present in cannabis as well.

Quality Control

• The quality assurance department report directly to the CEO, independent of production, marketing or other organization groups within the manufacturing company.
• The quality assurance procedures are revised on a periodic basis.
• The quality assurance inspection group has education, training, experience and understanding of their function.

Recall Capabilities

- **Initiation:** Recall is triggered upon identifying a product defect through customer complaints, production feedback, or instruction from the Ministry of Health (MOH).
- **Immediate Actions:** The QA Manager initiates the recall procedure, informs relevant regulatory authorities, and starts an investigation within 48 hours.
- **Customer Communication:** Defective products' recipients are identified and notified within 24 hours via email to ensure swift action and minimize risk.
- **Traceability and Accountability:** The Supply Chain Manager prepares detailed distribution records, while QA ensures the retrieval, segregation, and documentation of defective products.
- **Risk Management:** In consultation with the MOH, QA assesses the defect's impact and implements risk-reducing actions, including a public announcement if required.
- **Investigation Report:** A comprehensive investigation report, including root cause analysis, historical review, and corrective actions, is submitted to the MOH, with interim reports as needed.
- **Documentation and Closure:** QA maintains all recall documentation, ensures corrective actions to prevent recurrence, and reports recall efficiency to the MOH, ensuring full accountability and product safety.

Customer Complaints

- **Complaint Reception:** Complaints can be received via email, telephone, or other means. All complaints are documented immediately by the QA department within one working day.
- **Documentation:** The complaint is recorded with a unique consecutive number for traceability.
- **Investigation:** The QA department conducts a thorough investigation, including batch record reviews, risk assessments, and identification of the root cause, documented in the Customer Complaint - Investigation Report.
- **Customer Communication:** The QA department is responsible for correspondence with the customer, ensuring timely updates and final responses.
- **Corrective Actions:** Appropriate corrective and preventive actions (CAPA) are identified and implemented to eliminate recurring issues.

Customer Complaints

- **Product Status:** If the product's validity is in doubt, it may be quarantined. The QA Manager ensures no further distribution until the final status is determined.
- **Closure and Review:** Complaints are closed after completing all investigations and customer responses within 30 days. The QA Manager reviews complaint records regularly to identify trends and prevent future issues, reporting findings during the Product Quality Review and Management Review meetings.

3. Eybna's Compliance and Standards

We hereby declare that all of Eybna Technologies products fit the following criteria:

- Meet applicable requirements set out in the Food and Drug Regulations C.R.C., c. 870.
- Meet applicable requirements set out in the Safe Food for Canadians Regulations SOR/2018-108.
- Composed of raw materials that appear in GRAS list.
- Comply with Section 21 CFR of the Federal Food, Drug and Cosmetic Act.
- Comply with European Council Directive 88/388/EEC and EU Regulation 1334/2008.
- Certified OU Kosher.
- Composed of only natural ingredients as defined in Section 21 CFR 101.22.
- Meet the E.U. regulations as outlined in 1881/2006/EC in regards to contaminants prohibited in food grade raw material ingredients.
- Meet Proposed Requirements for Mandatory Testing of Pesticide Active Ingredients in Cannabis Products, Office of Medical Cannabis 144 (published in Canada Gazette, Part II, on July 11, 2018).
- Conforms to I-502 regulation, with regard to residual solvents.
- Compliant with Schedule B to the FDA.
- Compliant with: Regulation (EU) No 1169/2011, Regulation (EU) No 2023/915, Regulation (EC) No 1333/2008, Regulation (EC) No 1334/2008, Regulation (EU) 872/2012, Regulation (EU) 2018/73, Regulation (EC) No 2073/2005.

4. Shelf Life of Eybna's Products

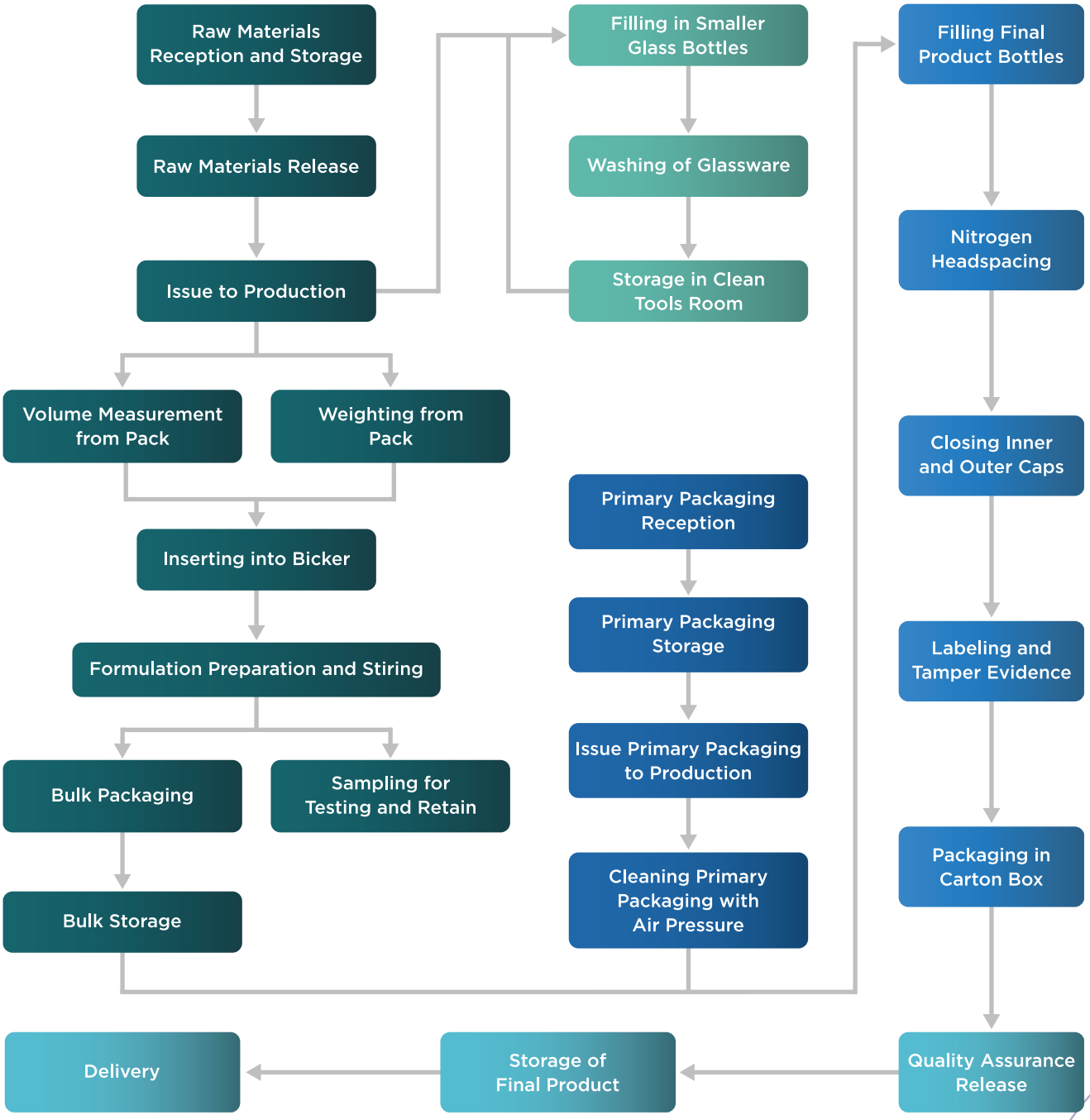
Eybna includes an expiration date for each batch on every label and in its supporting documents to signal how long the product is expected to preserve its intended terpene-flavour profile. This date marks optimal quality rather than a food-safety limit.

Because natural terpene blends can change over time, customers may notice gradual shifts in taste, colour, or pH—changes that advance faster or slower depending on storage conditions. To keep quality at its peak, store products as directed and protect them from oxygen, heat, light, moisture, and freezing.

After a container is opened, longevity depends on both the product's age and how well it is resealed and stored. Use opened formulations promptly; if they must be held, purge the headspace with nitrogen, close the lid tightly, and keep the product under the recommended conditions. Partially used containers should be finished as soon as feasible.

For best performance, apply Eybna formulations before the labelled expiration date. Products may still function after that point, but any post-date use should be based on the customer's own testing and is entirely at their own risk.

5. Eybna's Production FlowChart



6. Eybna's Process Description

Name	Description
Raw Material Reception	All the materials purchased are approved in Israel, based on EFSA risk assessment. From approved suppliers requires QA release that involves verification of COAs for the incoming batches + verification that the order matches the received materials + verification that the containers are unharmed and closed properly
Raw Material Storage	In RM warehouse in ambient temperature (hold and release sections are segregated in the warehouse)
Issue to Production	On stainless steel or plastic manual cart
Filling in Smaller Glass Bottles	Materials that used are frequently transmitted into multiple-use glass bottles, Under the hood, with a pump, intended specifically for terpene. The bottles pass washing, as described between different uses (after emptying). The bottles' completeness was checked every use. It is possible to add the same material from the same batch to the remaining material in the bottle, under the condition there is the same batch, not more than 2 months.
Washing of Glassware	With Ethanol and then drying in the oven. Bottle completeness checked after drying
Storage in Clean Tools Room	In ambient temperature, aerated, top-down, protected from breakage
Volume measurement from pack	Under hood with calibrated single use hard-plastic pipette
Weighing	Under hood w. calibrated analytical scale. Placed on a daily used plastic table with stainless steel tool washed after each use
Formulation Preparation - Inserting into stirer	Under hood. By hard plastic pipette or weighted, moved by stainless steel spoon. On the stirrer, there is a glass container inside which a magnet is used for stirring. Magnet cleaned after each use - by water with soap, food-grade ethanol, and then drying Magnet checked for its completeness every use
Stirring	Under hood. Time vary from 5 minutes to 120 minutes, speed - according to product recipe. The product is covered during mixing. The final release of the product includes visual inspection, density, RI, and odor, which ensures that mixing was enough
Sampling for Testing and Retain	Under hood. Done by clean **hard-plastic** pipette into 10ml aluminium bottles
Bulk Packaging	Under the hood - the glass/aluminum container is labeled with the product name, SKU, batch #, and manufacturing date. Head spacing with nitrogen and hermetical closing only for aluminum bottles
Bulk Storage	In production room - no requirement for cooling.

6. Eybna's Process Description

Name	Description
Bulk Storage	In production room - no requirement for cooling.
Primary Packaging Reception	From approved suppliers, Packs from glass and aluminium No release process - Chain supply verify the order matches the required items in the required quantity
Primary Packaging Storage	In ambient temperature in closed packs
Cleaning with air pressure	To verify no foreign matter is in the bottles. Air compressor with usage of oil. Air from compressor sampled annually for micro
Filling final product bottles	Under hood with single use hard plastic calibrated pipette or glass measure tool, washed after each use. Checked for completeness before each use
Nitrogen head spacing	Under hood, for a few seconds, during closure of cap. Performed for final product bottles and also raw material and bulk containers
Closing inner cap	Under hood while using nitrogen to fill the head space.
Closing outer cap	Under hood. No contact with the product
Labeling & tamper evidence	Labeling - for industry. Labels first version is checked according to regulations in country of use. Includes verification that the correct labels were used. Tamper evidence by sticking from cap to bottle
Carron box producing	Folding the carton
Packing in carton box	Manual Placing labeled bottle w. tamper evidence into the box and closing.
Carton labeling	Same label as on bottle
Quality Assurance Final release	QA checks the bottles (verification that the product is undamaged, properly closed, w. required packaging materials, label data conforms to required details, and all required tests such as Density, RI, Odor, and Visual. Additional tests may be done according to customer requirements. Additionally, the products are sampled for pesticides, heavy metals, and residual solvents according to the sampling plan
Storage of Final Product	In final product warehouse in ambient temperature
Delivery	Packaging per SO - with proper labeling and pictograms according to product SDS

7. Recall Plan

1. **Purpose:** The Recall Procedure ensures that any product that is deemed unsafe or non-compliant with regulatory standards can be quickly and effectively removed from the market to protect public health and maintain regulatory compliance.
2. **Scope:** This procedure applies to all products manufactured, packaged, or distributed by the company. It includes any product that has been identified as potentially harmful or non-compliant.
3. **Recall Process:**

Identification and Evaluation:

- Identify the need for a recall through internal testing, customer complaints, or regulatory notification.
- Evaluate the severity and scope of the issue to determine the recall classification.

Initiation:

- Notify the Recall Coordinator immediately.
- Form a recall team including QA, Regulatory Affairs, and other relevant departments.

Communication:

- Notify regulatory authorities and provide them with the necessary information about the recall.
- Inform customers, distributors, and the public through appropriate channels (e.g., press releases, direct communication).

Execution:

- Retrieve affected products from the market.
- Implement measures to prevent further distribution or sale.
- Conduct a root cause analysis to understand the reason for the defect or non-compliance.

Disposition:

- Decide the appropriate disposition of the recalled products (e.g., destruction, rework).
- Document all actions taken during the recall process.

Follow-up and Effectiveness Checks:

- Verify that all affected products have been recalled.
- Evaluate the effectiveness of the recall process and make improvements as necessary.

Documentation:

- Maintain detailed records of the recall process, including communications, actions taken, and final disposition of the products.
- Ensure all documentation is readily available for regulatory review.

8. Food-GMP Certification and Pharmaceutical Compatibility of Terpene-Based Formulations

Flavours are routinely used as pharmaceutical excipients to improve palatability. Although full pharmaceutical GMP (ICH Q7) is more stringent than food GMP, both systems share the same quality pillars: documented procedures, validated test methods, traceability, and change control. Eybna now holds both FSSC 22000 certification and a Good Manufacturing Practice (GMP) certificate issued by the Food Division of the Israeli Ministry of Health (IMoH). These credentials confirm that our documented quality system governs the entire life-cycle—design, production, research, and distribution—of every terpene-based formulation we manufacture.

Key points for pharmaceutical customers:

- Documented Quality System – All processes are controlled by written SOPs, batch records, risk-based change control, and full traceability; documents can be shared under NDA.
- Raw-Material Standards – Formulations use FEMA-GRAS substances and natural botanical isolates that meet the purity criteria of EU Regulation No. 872/2012 and comply with the compositional and labelling requirements of EU Regulation No. 1334/2008 on flavourings.
- Regulatory Alignment – The International Pharmaceutical Excipients Council (IPEC) recognises that FEMA-GRAS safety evaluations may be referenced in a Drug Master File (DMF) to support excipient use.
- Scope of Certification – Our GMP certificate is issued by a food-regulatory authority. While it demonstrates strong GMP compliance, it does not by itself constitute formal pharmaceutical GMP certification; additional qualification may be required by individual drug manufacturers or health authorities.

Taken together, FSSC 22000, IMoH Food-GMP certification, and conformity with EU 872/2012 and EU 1334/2008 provide a robust quality platform that supports the use of Eybna terpene formulations as excipients in many pharmaceutical applications. Customers who require full pharmaceutical GMP alignment are welcome to review our quality documentation and perform on-site audits as needed.

9. Allergens Statement

We hereby confirm that the following allergens are not present in Eybna's Terpenes Formulations and that the manufacturer's procedure prevents allergen cross contamination in products.

Allergens	Present In The Products	Present in other products manufactured on the same line
Peanuts and peanut products	□ Yes ■ No	□ Yes ■ No
Milk and milk products	□ Yes ■ No	□ Yes ■ No
Eggs and egg products	□ Yes ■ No	□ Yes ■ No
Fish and fish products	□ Yes ■ No	□ Yes ■ No
Shellfish	□ Yes ■ No	□ Yes ■ No
Soy	□ Yes ■ No	□ Yes ■ No
Wheat and wheat products	□ Yes ■ No	□ Yes ■ No
Sulphites	□ Yes ■ No	□ Yes ■ No
Tree Nuts	□ Yes ■ No	□ Yes ■ No
Sesame Seeds	□ Yes ■ No	□ Yes ■ No
Mustard	□ Yes ■ No	□ Yes ■ No

Notice:
This Allergens Checklist does not relieve the purchaser from undertaking their own tests in order to assure the above-mentioned allergens are not present in the product.

Gil Tsapovetsky
Quality Assurance Manager

10. Genetically Modified Organisms (GMO) Statement

Genetically Modified Organisms (GMO)	
Are Eybna's Terpenes Formulations or any of its ingredients derived from any genetically modified materials?	□ Yes ■ No
Do the final products, or any of its ingredients contain any genetically modified materials?	□ Yes ■ No
Were Eybna's Terpenes Formulations products irradiated or contain any irradiated ingredients?	□ Yes ■ No
Do our products require labeling under EU Regulation 1829/2003 and 1820/2003?	□ Yes ■ No
Do the products consist of GMOs, contain GMOs, or is it produced from GMOs as defined in the EU Regulation 1829/2003 and 1830/2003?	□ Yes ■ No

Notice:
This GMO Questionnaire does not relieve the purchaser from undertaking their own tests in order to assure the suitability of this product for its application and to comply with all relevant legal requirements for any goods into which this product may be in.

Gil Tsapovetsky
Quality Assurance Manager

11. Certificate of Origin

Supplier Name: Eybna Technologies Ltd.

We hereby declare that Eybna's Terpenes Formulations were produced by Eybna Technologies Ltd. In Israel.

Gil Tsapovetsky
Quality Assurance Manager

12. Food Grade Certificate

This is to verify that all the ingredients of our products purchased from Eybna Technologies Ltd. are food grade quality.

The materials are allowed for use as flavor ingredients intended for human consumption, or as indirect additives according to Section 21 CFR of the Federal Food, Drug and Cosmetic Act.

They do also comply with the European Council Directive EU Regulation 1334/2008.

Gil Tsapovetsky
Quality Assurance Manager

13. Natural Certificate

We are pleased to certify to you that at the time of shipment, Eybna's Terpenes Formulations are on such date not adulterated or misbranded within the meaning of the Federal Food, Drug and Cosmetic Act.

The product is not an article which may not, under the provisions of Section 404 or 405 of said act be introduced into interstate commerce.

Eybna's Terpenes Formulations are not adulterated or misbranded within the meaning of the food or drug laws of any state or municipality which are applicable to such shipment or delivery. The product is composed of only natural ingredients as defined in Section 21 CFR 101.22 and to the best of our knowledge and belief contains no artificial flavor ingredients.

The guarantees given herein are continuing and shall be in full force and effect until revoked in writing. Please contact us at your convenience in case of requiring additional information.

Notice:
This Natural Certificate does not relieve the purchaser from undertaking their own tests in order to assure the suitability of this product for its application and to comply with all relevant legal requirements for any goods into which this product may be in.

Gil Tsapovetsky
Quality Assurance Manager

14. Shelf Life Statement

Product Name: Pineapple Haze

Product Code: 8-02-9999

We hereby declare that the product has a shelf life of 24 months from date of production.

Eybna Technologies recommends that the goods will be stored in their original bottles in an ambient temperature. It is recommended to prevent repetitive reopening of the bottles.

In the case of reopening, it is recommended to reseal bottles with nitrogen gas.

Gil Tsapovetsky
Quality Assurance Manager

15. Sanitary Conditions Statement

Supplier Name: Eybna Technologies Ltd.

We hereby declare that all products produced by Eybna Technologies Ltd. are manufactured, prepared, preserved, packaged, and stored under strict sanitary conditions to prevent microbial contamination.

Gil Tsapovetsky
Quality Assurance Manager

16. GRAS Declaration

Supplier Name: Eybna Technologies Ltd.
Product Name: Kush Note
Product Code: 7-500-9999

This is to declare that the above item, purchased from EybnaTechnologies is composed of only GRAS ingredients that were approved by the FDA for usein flavors under 21 CF.

Gil Tsapovetsky
Quality Assurance Manager

17. Product Composition Statement

Date of Print: 7. August 24
Supplier Name: Eybna Technologies Ltd.
Kush Note7-500-9999

We hereby declare that that the above-mentioned product contains only botanical -derived compounds and that the following materials are not being used to manufacture the above product:

- Squalene
- Squalane
- Amino acids
- Caffeine
- Colouring agents
- Essential fatty acids
- Glucuronolactone
- Probiotics
- Taurine
- Vitamins
- Mineral nutrients
- Diacetyl
- 2,3-pentanedione
- Preservatives
- (PAHs) polycyclic aromatic hydrocarbons
- Naphtalene
- 1,3-Butandiene
- Propylene Oxide
- Benzene
- Toluene
- Nitrosamines

Notice: This document does not relieve the purchaser from undertaking their own tests in order to assure the suitability of this product for its application and to comply with all relevant legal requirements for any goods into which this product may be in.

Gil Tsapovetsky
Quality Assurance Manager

18. Non-Cannabis Goods Statement

This certificate is issued to confirm that the product specified below is free from any cannabis or hemp-derived components.

- Product Name: Kush Note - Enhancer Line
- SKU: 7-500-9999

We hereby certify that the above-mentioned product, [Product Name], does not contain any cannabis or hemp-derived ingredients. Our stringent quality control processes ensure that this product is composed solely of high-quality, all-natural ingredients, providing a pure and unadulterated flavor experience.

This declaration is made with the understanding that it assures customers and regulatory bodies of the non-cannabis composition of the product.

Gil Tsapovetsky
Quality Assurance Manager

19. BSE/TSE Statement

Product Name: Kush Note - Enhancer Line **Product Code: 7-500-9999**

The following product is composed of natural, botanically derived compounds and does not contain any animal products or any materials of animal origin. The product is free from BSE (Bovine Spongiform Encephalopathy) and TSE (Transmissible Spongiform Encephalopathy).

Gil Tsapovetsky
Quality Assurance Manager

20. Irradiation Statement

Product Name: Kush Note - Enhancer Line **Product Code: 7-500-9999**

We hereby declare that our product “Invigorate” and the materials used in its production have not been subject to irradiation of any kind during their manufacture.

Gil Tsapovetsky
Quality Assurance Manager

21. Technical Data Sheet

Technical Data Sheet

Date of Print: 4. June 25
Product Name: Kush Note - Enhancer Line
Product Code: 7-500-9999

Natural status: This product is natural according to Section 21 CFR 101.22.

GMO status: This product contains no genetically modified materials, according to the EU Regulations 1829/2003, 1830/2003 and 1820/2003.

Additives: This product contains no additives.

Food allergens: This product is free from recognized food allergens as defined by EU Regulation 1169/2011 and FALCPA.

Kosher status: This product is certified kosher and suitable for kosher use.

Halal status: This product is suitable for halal use, but not certified as such.

Food grade status: This product is assembled only of materials allowed for use as flavor ingredients intended for human consumption, or as indirect additives according to Section 21 CFR of the Federal Food, Drug and Cosmetic Act.

Organic Compliance: This product only contains permitted ingredients that are on the National List of non-organic ingredients permitted in certified organic agriculture and processing.

Packaging Sizes: This product is available in 10ml, 50ml, 250ml, and 1.25 Liter aluminium bottles. Each size is designed to ensure optimal usage and storage efficiency.

Storage: Keep container tightly closed in a dry and well-ventilated place. Containers which are opened must be carefully resealed and kept upright to prevent leakage.

Product Use: This product is only for industrial use to be added to final products and not for direct consumption.

Notice:
This Technical Data Sheet does not relieve the purchaser from undertaking their own tests in order to assure the suitability of this product for its application and to comply with all relevant legal requirements for any goodsinto which this product may be in.

Gil Tsapovetksy
QA Manager

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22. Safety Data Sheet

Safety Data Sheet

acc. to 29 CFR 1910.1200 App D

Kush Note

Version number: GHS 1.0 Date of compilation: 2024-05-26

SECTION 1: Identification

1.1 Product identifier

Trade name **Kush Note**

Product number 7-500-9999

1.2 Relevant identified uses of the substance or mixture and uses advised against

Relevant identified uses Industrial use

1.3 Details of the supplier of the safety data sheet

Eybna Technologies Ltd
1 HaYozma St.
4442214 Kfar Saba
Israel
Telephone: +972 3 3741976
e-mail: info@eybna.com
Website: http://www.eybna.com/

e-mail (competent person) Gil.ts@eybna.com (Gil Tsapovetsky)

1.4 Emergency telephone number +1 4158544820

SECTION 2: Hazard(s) identification

2.1 Classification of the substance or mixture

Classification acc. to OSHA "Hazard Communication Standard" (29 CFR 1910.1200)

Section	Hazard class	Category	Hazard class and category	Hazard statement
A.1O	acute toxicity (oral)	4	Acute Tox. 4	H302
A.2	skin corrosion/irritation	2	Skin Irrit. 2	H315
A.3	serious eye damage/eye irritation	2	Eye Irrit. 2	H319
A.4S	skin sensitization	1	Skin Sens. 1	H317
A.6	carcinogenicity	2	Carc. 2	H351
A.10	aspiration hazard	1	Asp. Tox. 1	H304
B.6	flammable liquid	3	Flam. Liq. 3	H226

For full text of abbreviations: see SECTION 16.

The most important adverse physicochemical, human health and environmental effects

The product is combustible and can be ignited by potential ignition sources.

2.2 Label elements

Labelling acc. to OSHA "Hazard Communication Standard" (29 CFR 1910.1200)

- Signal word danger
- Pictograms

GHS02, GHS07, GHS08 

United States: en Page: 1 / 15

22. Safety Data Sheet

- Hazard statements

H226	Flammable liquid and vapor.
H302	Harmful if swallowed.
H304	May be fatal if swallowed and enters airways.
H315	Causes skin irritation.
H317	May cause an allergic skin reaction.
H319	Causes serious eye irritation.
H351	Suspected of causing cancer.

- Precautionary statements

P202	Do not handle until all safety precautions have been read and understood.
P210	Keep away from heat/sparks/open flames/hot surfaces. No smoking.
P233	Keep container tightly closed.
P240	Ground/bond container and receiving equipment.
P241	Use explosion-proof electrical/ventilating/lighting equipment.
P242	Use only non-sparking tools.
P243	Take precautionary measures against static discharge.
P261	Avoid breathing dust/fume/gas/mist/vapors/spray.
P270	Do not eat, drink or smoke when using this product.
P272	Contaminated work clothing must not be allowed out of the workplace.
P280	Wear protective gloves/eye protection/face protection.
P301+P310	If swallowed: Immediately call a poison center/doctor.
P301+P312	If swallowed: Call a poison center/doctor if you feel unwell.
P302+P352	If on skin: Wash with plenty of water.
P303+P361+P353	If on skin (or hair): Take off immediately all contaminated clothing. Rinse skin with water/ shower.
P305+P351+P338	If in eyes: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing.
P308+P313	If exposed or concerned: Get medical advice/attention.
P321	Specific treatment (see on this label).
P330	Rinse mouth.
P331	Do NOT induce vomiting.
P333+P313	If skin irritation or rash occurs: Get medical advice/attention.
P337+P313	If eye irritation persists: Get medical advice/attention.
P362	Take off contaminated clothing and wash before reuse.
P363	Wash contaminated clothing before reuse.
P370+P378	In case of fire: Use sand, carbon dioxide or powder extinguisher to extinguish.
P403+P235	Store in a well-ventilated place. Keep cool.
P405	Store locked up.
P501	Dispose of contents/container to industrial combustion plant.

2.3 Other hazards

Hazards not otherwise classified

Very toxic to aquatic life with long lasting effects (GHS category 1: aquatic toxicity - acute and/or chronic).

Results of PBT and vPvB assessment

Does not contain a PBT-/vPvB-substance at a concentration of ≥ 0.1%.

Endocrine disrupting properties

Does not contain an endocrine disruptor (ED) in a concentration of ≥ 0.1%.

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22. Safety Data Sheet

SECTION 3: Composition/information on ingredients

3.1 Substances

Not relevant (mixture)

3.2 Mixtures

Description of the mixture

Name of substance	Wt%	Classification acc. to GHS
Proprietary Monoterpene	25 – < 50	Skin Irrit. 2 / H315 Eye Irrit. 2 / H319 Carc. 2 / H351 Asp. Tox. 1 / H304 Flam. Liq. 3 / H226
Proprietary Sesquiterpene	25 – < 50	Acute Tox. 4 / H302
Proprietary Monoterpene	1 – < 5	Acute Tox. 4 / H302 Skin Irrit. 2 / H315 Eye Irrit. 2 / H319 Skin Sens. 1 / H317 STOT SE 3 / H335 Flam. Liq. 3 / H226
Proprietary Monoterpenic Alcohol	1 – < 5	Skin Irrit. 2 / H315 Eye Irrit. 2 / H319 STOT SE 3 / H335
Proprietary Sesquiterpene	1 – < 5	Acute Tox. 4 / H302

Remarks

For full text of abbreviations: see SECTION 16

SECTION 4: First-aid measures

4.1 Description of first-aid measures

General notes

Do not leave affected person unattended. Remove victim out of the danger area. Keep affected person warm, still and covered. Take off immediately all contaminated clothing. In all cases of doubt, or when symptoms persist, seek medical advice. In case of unconsciousness place person in the recovery position. Never give anything by mouth.

Following inhalation

If breathing is irregular or stopped, immediately seek medical assistance and start first aid actions. In case of respiratory tract irritation, consult a physician. Provide fresh air.

Following skin contact

Wash with plenty of soap and water.

Following eye contact

Remove contact lenses, if present and easy to do. Continue rinsing. Irrigate copiously with clean, fresh water for at least 10 minutes, holding the eyelids apart.

Following ingestion

Rinse mouth with water (only if the person is conscious). Do NOT induce vomiting.

4.2 Most important symptoms and effects, both acute and delayed

Symptoms and effects are not known to date.

4.3 Indication of any immediate medical attention and special treatment needed

none

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22. Safety Data Sheet

Safety Data Sheet

acc. to 29 CFR 1910.1200 App D

Kush Note

Version number: GHS 1.0 Date of compilation: 2024-05-26

SECTION 5: Fire-fighting measures

5.1 Extinguishing media

Suitable extinguishing media

Water spray, BC-powder, Carbon dioxide (CO2)

Unsuitable extinguishing media

Water jet

5.2 Special hazards arising from the substance or mixture

In case of insufficient ventilation and/or in use, may form flammable/explosive vapor-air mixture. Solvent vapors are heavier than air and may spread along floors. Places which are not ventilated, e.g. unventilated below ground level areas such as trenches, conduits and shafts, are particularly prone to the presence of flammable substances or mixtures.

Hazardous combustion products

Carbon monoxide (CO), Carbon dioxide (CO2)

5.3 Advice for firefighters

In case of fire and/or explosion do not breathe fumes. Coordinate firefighting measures to the fire surroundings. Do not allow firefighting water to enter drains or water courses. Collect contaminated firefighting water separately. Fight fire with normal precautions from a reasonable distance.

SECTION 6: Accidental release measures

6.1 Personal precautions, protective equipment and emergency procedures

For non-emergency personnel

Remove persons to safety.

For emergency responders

Wear breathing apparatus if exposed to vapors/dust/aerosols/gases.

6.2 Environmental precautions

Keep away from drains, surface and ground water. Retain contaminated washing water and dispose of it. If substance has entered a water course or sewer, inform the responsible authority.

6.3 Methods and material for containment and cleaning up

Advice on how to contain a spill

Covering of drains

Advice on how to clean up a spill

Wipe up with absorbent material (e.g. cloth, fleece). Collect spillage: sawdust, kieselgur (diatomite), sand, universal binder

Appropriate containment techniques

Use of adsorbent materials.

Other information relating to spills and releases

Place in appropriate containers for disposal. Ventilate affected area.

6.4 Reference to other sections

Hazardous combustion products: see section 5. Personal protective equipment: see section 8. Incompatible materials: see section 10. Disposal considerations: see section 13.

22. Safety Data Sheet

SECTION 7: Handling and storage

7.1 Precautions for safe handling

Recommendations

- Measures to prevent fire as well as aerosol and dust generation

Use local and general ventilation. Avoidance of ignition sources. Keep away from sources of ignition - No smoking. Take precautionary measures against static discharge. Use only in well-ventilated areas. Due to danger of explosion, prevent leakage of vapours into cellars, flues and ditches. Ground/bond container and receiving equipment. Use explosion-proof electrical/ventilating/lighting/equipment. Use only non-sparking tools.

- Specific notes/details

Places which are not ventilated, e.g. unventilated below ground level areas such as trenches, conduits and shafts, are particularly prone to the presence of flammable substances or mixtures. Vapors are heavier than air, spread along floors and form explosive mixtures with air. Vapors may form explosive mixtures with air.

Advice on general occupational hygiene

Wash hands after use. Do not eat, drink and smoke in work areas. Remove contaminated clothing and protective equipment before entering eating areas. Never keep food or drink in the vicinity of chemicals. Never place chemicals in containers that are normally used for food or drink. Keep away from food, drink and animal feedingstuffs.

7.2 Conditions for safe storage, including any incompatibilities

Managing of associated risks

- Explosive atmospheres

Keep container tightly closed and in a well-ventilated place. Use local and general ventilation. Keep cool. Protect from sunlight.

- Flammability hazards

Keep away from sources of ignition - No smoking. Keep away from heat, hot surfaces, sparks, open flames and other ignition sources. No smoking. Take precautionary measures against static discharge. Protect from sunlight.

- Ventilation requirements

Use local and general ventilation. Ground/bond container and receiving equipment.

- Packaging compatibilities

Only packagings which are approved (e.g. acc. to the Dangerous Goods Regulations) may be used.

7.3 Specific end use(s)

See section 16 for a general overview.

SECTION 8: Exposure controls/personal protection

8.1 Control parameters

Occupational exposure limit values (Workplace Exposure Limits)

Country	Name of substance	Identifier	TWA [ppm]	TWA [mg/m³]	STEL [ppm]	STEL [mg/m³]	Ceiling-C [ppm]	Ceiling-C [mg/m³]	Notation	Source
US		TLV®	20							ACGIH® 2019

Notation

Ceiling-C	ceiling value is a limit value above which exposure should not occur
STEL	short-term exposure limit: a limit value above which exposure should not occur and which is related to a 15-minute period (unless otherwise specified)
TWA	time-weighted average (long-term exposure limit): measured or calculated in relation to a reference period of 8 hours time-weighted average (unless otherwise specified

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22. Safety Data Sheet

Relevant DNELs of components

Endpoint	Threshold level	Protection goal, route of exposure	Used in	Exposure time
DNEL	9.03 mg/m³	human, inhalatory	worker (industry)	chronic - systemic effects
DNEL	158 mg/kg bw/day	human, dermal	worker (industry)	chronic - systemic effects
DNEL	3.8 mg/m³	human, inhalatory	worker (industry)	chronic - systemic effects
DNEL	0.542 mg/kg bw/day	human, dermal	worker (industry)	chronic - systemic effects

Relevant PNECs of components

Other names or synonyms	Endpoint	Threshold level	Organism	Environmental compartment	Exposure time
Proprietary Monoterpenic Alcohol	PNEC	68 µg/l	aquatic organisms	freshwater	short-term (single instance)
Proprietary Monoterpenic Alcohol	PNEC	6.8 µg/l	aquatic organisms	marine water	short-term (single instance)
Proprietary Monoterpenic Alcohol	PNEC	2.6 mg/l	aquatic organisms	sewage treatment plant (STP)	short-term (single instance)
Proprietary Monoterpenic Alcohol	PNEC	1.85 mg/kg	aquatic organisms	freshwater sediment	short-term (single instance)
Proprietary Monoterpenic Alcohol	PNEC	0.185 mg/kg	aquatic organisms	marine sediment	short-term (single instance)
Proprietary Monoterpenic Alcohol	PNEC	0.329 mg/kg	terrestrial organisms	soil	short-term (single instance)
Proprietary Monoterpene	PNEC	0.606 µg/l	aquatic organisms	freshwater	short-term (single instance)
Proprietary Monoterpene	PNEC	0.061 µg/l	aquatic organisms	marine water	short-term (single instance)
Proprietary Monoterpene	PNEC	0.2 mg/l	aquatic organisms	sewage treatment plant (STP)	short-term (single instance)
Proprietary Monoterpene	PNEC	157 µg/kg	aquatic organisms	freshwater sediment	short-term (single instance)
Proprietary Monoterpene	PNEC	15.7 µg/kg	aquatic organisms	marine sediment	short-term (single instance)
Proprietary Monoterpene	PNEC	31.7 µg/kg	terrestrial organisms	soil	short-term (single instance)

8.2 Exposure controls

Appropriate engineering controls

General ventilation.

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22. Safety Data Sheet

Individual protection measures (personal protective equipment)

Eye/face protection

Wear eye/face protection.

Skin protection

- Hand protection

Wear suitable gloves. Chemical protection gloves are suitable, which are tested according to EN 374. Check leak-tightness/impermeability prior to use. In the case of wanting to use the gloves again, clean them before taking off and air them well. For special purposes, it is recommended to check the resistance to chemicals of the protective gloves mentioned above together with the supplier of these gloves.

- Other protection measures

Take recovery periods for skin regeneration. Preventive skin protection (barrier creams/ointments) is recommended. Wash hands thoroughly after handling.

Respiratory protection

In case of inadequate ventilation wear respiratory protection.

Environmental exposure controls

Use appropriate container to avoid environmental contamination. Keep away from drains, surface and ground water.

SECTION 9: Physical and chemical properties

9.1 Information on basic physical and chemical properties

Appearance

Physical state	liquid
Color	
Particle	not relevant (liquid)
Odor	characteristic

Other safety parameters

pH (value)	not determined
Melting point/freezing point	not determined
Initial boiling point and boiling range	154.3 °C at 1,010 hPa
Flash point	31 °C at 1 atm
Evaporation rate	Not determined
Flammability (solid, gas)	not relevant, (fluid)
Vapor pressure	690 Pa at 20 °C
Density	not determined
Vapor density	this information is not available
Relative density	Information on this property is not available

22. Safety Data Sheet

	Solubility(ies)	not determined

Partition coefficient

- n-octanol/water (log KOW)	this information is not available

Auto-ignition temperature	245 °C
Viscosity	not determined
Explosive properties	none
Oxidizing properties	none

9.2	**Other information**	there is no additional information

SECTION 10: Stability and reactivity

10.1 Reactivity

Concerning incompatibility: see below "Conditions to avoid" and "Incompatible materials". The mixture contains reactive substance(s). Risk of ignition.

If heated:

Risk of ignition

10.2 Chemical stability

See below "Conditions to avoid".

10.3 Possibility of hazardous reactions

No known hazardous reactions.

10.4 Conditions to avoid

Keep away from heat, hot surfaces, sparks, open flames and other ignition sources. No smoking.

Hints to prevent fire or explosion

Use explosion-proof electrical/ventilating/lighting/equipment. Use only non-sparking tools. Take precautionary measures against static discharge.

10.5 Incompatible materials

Oxidizers

10.6 Hazardous decomposition products

Reasonably anticipated hazardous decomposition products produced as a result of use, storage, spill and heating are not known. Hazardous combustion products: see section 5.

SECTION 11: Toxicological information

11.1 Information on toxicological effects

Test data are not available for the complete mixture.

Classification procedure

The method for classification of the mixture is based on ingredients of the mixture (additivity formula).

Classification acc. to OSHA "Hazard Communication Standard" (29 CFR 1910.1200)

Acute toxicity

Harmful if swallowed.

22. Safety Data Sheet

- Acute toxicity estimate (ATE)
  Oral 1,002 $^{mg}/_{kg}$

Acute toxicity estimate (ATE) of components		
Other names or synonyms	**Exposure route**	**ATE**
Proprietary Sesquiterpene	oral	500 $^{mg}/_{kg}$
Proprietary Sesquiterpene	oral	500 $^{mg}/_{kg}$
Proprietary Monoterpene	oral	500 $^{mg}/_{kg}$

Skin corrosion/irritation
Causes skin irritation.

Serious eye damage/eye irritation
Causes serious eye irritation.

Respiratory or skin sensitization
May cause an allergic skin reaction.

Germ cell mutagenicity
Shall not be classified as germ cell mutagenic.

Carcinogenicity
Suspected of causing cancer.

IARC Monographs on the Evaluation of Carcinogenic Risks to Humans		
Name of substance	**Classification**	**Number**
Proprietary Monoterpene	2B	

Legend
2B Possibly carcinogenic to humans

Reproductive toxicity
Shall not be classified as a reproductive toxicant.

Specific target organ toxicity - single exposure
Shall not be classified as a specific target organ toxicant (single exposure).

Specific target organ toxicity - repeated exposure
Shall not be classified as a specific target organ toxicant (repeated exposure).

Aspiration hazard
May be fatal if swallowed and enters airways.

SECTION 12: Ecological information

12.1 Toxicity
Very toxic to aquatic life with long lasting effects.

22. Safety Data Sheet

Aquatic toxicity (acute) of components

Other names or synonyms	Endpoint	Value	Species	Exposure time
Proprietary Monoterpene	EC50	1.47 $^{mg}/_{l}$	aquatic invertebrates	48 h
Proprietary Monoterpene	ErC50	0.342 $^{mg}/_{l}$	algae	72 h
Proprietary Monoterpenic Alcohol	LC50	>82 $^{mg}/_{l}$	fish	96 h
Proprietary Monoterpenic Alcohol	EC50	10 $^{mg}/_{l}$	aquatic invertebrates	48 h
Proprietary Monoterpenic Alcohol	ErC50	>11 $^{µg}/_{l}$	algae	72 h
Proprietary Monoterpene	LC50	0.303 $^{mg}/_{l}$	fish	96 h
Proprietary Monoterpene	EC50	0.475 $^{mg}/_{l}$	aquatic invertebrates	48 h

12.2 Persistence and degradability

Data are not available.

12.3 Bioaccumulative potential

Data are not available.

12.4 Mobility in soil

Data are not available.

12.5 Results of PBT and vPvB assessment

According to the results of its assessment, this substance is not a PBT or a vPvB. Does not contain a PBT-/vPvB-substance at a concentration of ≥ 0.1%.

12.6 Endocrine disrupting properties

Does not contain an endocrine disruptor (ED) in a concentration of ≥ 0.1%.

12.7 Other adverse effects

Data are not available.

SECTION 13: Disposal considerations

13.1 Waste treatment methods

Waste treatment-relevant information

Solvent reclamation/regeneration.

Sewage disposal-relevant information

Do not empty into drains. Avoid release to the environment. Refer to special instructions/safety data sheets.

Waste treatment of containers/packages

Only packagings which are approved (e.g. acc. to DOT) may be used. Completely emptied packages can be recycled. Handle contaminated packages in the same way as the substance itself.

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22. Safety Data Sheet

Remarks

Please consider the relevant national or regional provisions. Waste shall be separated into the categories that can be handled separately by the local or national waste management facilities.

SECTION 14: Transport information

14.1 UN number

DOT	UN 1993
IMDG-Code	UN 1993
ICAO-TI	UN 1993

14.2 UN proper shipping name

DOT	Flammable liquid, n.o.s.
IMDG-Code	FLAMMABLE LIQUID, N.O.S.
ICAO-TI	Flammable liquid, n.o.s.
Technical name (hazardous ingredients)	Myrcene, Alpha-Pinene

14.3 Transport hazard class(es)

DOT	3
IMDG-Code	3
ICAO-TI	3

14.4 Packing group

DOT	III
IMDG-Code	III
ICAO-TI	III

14.5 Environmental hazards hazardous to the aquatic environment

Environmentally hazardous substance (aquatic environment) Myrcene

14.6 Special precautions for user

There is no additional information.

14.7 Transport in bulk according to IMO instruments

The cargo is not intended to be carried in bulk.

Information for each of the UN Model Regulations

Transport of dangerous goods by road or rail (49 CFR US DOT) - Additional information

Particulars in the shipper's declaration	UN1993, Flammable liquid, n.o.s., (contains: Myrcene, Alpha-Pinene), 3, III, environmentally hazardous
Danger label(s)	3, fish and tree

Environmental hazards	yes (hazardous to the aquatic environment)
Special provisions (SP)	B1, B52, IB3, T4, TP1, TP29
ERG No	128

22. Safety Data Sheet

International Maritime Dangerous Goods Code (IMDG) - Additional information

Marine pollutant	yes (hazardous to the aquatic environment) (Myrcene)
Danger label(s)	3, fish and tree

Special provisions (SP)	223, 274, 955
Excepted quantities (EQ)	E1
Limited quantities (LQ)	5 L
EmS	F-E, S-E
Stowage category	A

International Civil Aviation Organization (ICAO-IATA/DGR) - Additional information

Environmental hazards	yes (hazardous to the aquatic environment)
Danger label(s)	3

Special provisions (SP)	A3
Excepted quantities (EQ)	E1
Limited quantities (LQ)	10 L

SECTION 15: Regulatory information

15.1 Safety, health and environmental regulations specific for the product in question

National regulations (United States)

Toxic Substance Control Act (TSCA) not all ingredients are listed (ACTIVE)

Superfund Amendment and Reauthorization Act (SARA TITLE III)

- The List of Extremely Hazardous Substances and Their Threshold Planning Quantities (EPCRA Section 302, 304)

none of the ingredients are listed

- Specific Toxic Chemical Listings (EPCRA Section 313)

none of the ingredients are listed

Comprehensive Environmental Response, Compensation, and Liability Act (CERCLA)

- List of Hazardous Substances and Reportable Quantities (CERCLA section 102a) (40 CFR 302.4)

none of the ingredients are listed

Clean Air Act

none of the ingredients are listed

Right to Know Hazardous Substance List

- Hazardous Substance List (NJ-RTK)

Name of substance	CAS No	Remarks	Classifications
Alpha-Pinene	80-56-8		F3

Legend

F3 Flammable - Third Degree

22. Safety Data Sheet

California Environmental Protection Agency (Cal/EPA): Proposition 65 - Safe Drinking Water and Toxic Enforcement Act of 1987

Proposition 65 List of chemicals

Name acc. to inventory	CAS No	Remarks	Type of the toxicity
beta-Myrcene	123-35-3		cancer

Industry or sector specific available guidance(s)

NPCA-HMIS® III

Hazardous Materials Identification System. American Coatings Association.

Category	Rating	Description
Chronic	*	chronic (long-term) health effects may result from repeated overexposure
Health	2	temporary or minor injury may occur
Flammability	3	material that can be ignited under almost all ambient temperature conditions
Physical hazard	0	material that is normally stable, even under fire conditions, and will not react with water, polymerize, decompose, condense, or self-react. Non-explosive
Personal protection	-	

NFPA® 704

National Fire Protection Association: Standard System for the Identification of the Hazards of Materials for Emergency Response (United States).

Category	Degree of hazard	Description
Flammability	3	material that can be ignited under almost all ambient temperature conditions
Health	2	material that, under emergency conditions, can cause temporary incapacitation or residual injury
Instability	0	material that is normally stable, even under fire conditions
Special hazard		

National inventories

Country	Inventory	Status
US	TSCA	not all ingredients are listed

Legend
TSCA Toxic Substance Control Act

15.2 Chemical Safety Assessment

Chemical safety assessments for substances in this mixture were not carried out.

22. Safety Data Sheet

SECTION 16: Other information, including date of preparation or last revision

Abbreviations and acronyms

Abbr.	Descriptions of used abbreviations
49 CFR US DOT	49 CFR U.S. Department of Transportation
ACGIH® 2019	From ACGIH®, 2019 TLVs® and BEIs® Book. Copyright 2019. Reprinted with permission. Information on the proper use of the TLVs® and BEIs®: http://www.acgih.org/tlv-bei-guidelines/policies-procedures-presentations/tlv-bei-position-statement
Acute Tox.	Acute toxicity
Asp. Tox.	Aspiration hazard
ATE	Acute Toxicity Estimate
Carc.	Carcinogenicity
CAS	Chemical Abstracts Service (service that maintains the most comprehensive list of chemical substances)
Ceiling-C	Ceiling value
DGR	Dangerous Goods Regulations (see IATA/DGR)
DNEL	Derived No-Effect Level
DOT	Department of Transportation (USA)
EC50	Effective Concentration 50 %. The EC50 corresponds to the concentration of a tested substance causing 50 % changes in response (e.g. on growth) during a specified time interval
ED	Endocrine disruptor
EmS	Emergency Schedule
ErC50	≡ EC50: in this method, that concentration of test substance which results in a 50 % reduction in either growth (EbC50) or growth rate (ErC50) relative to the control
ERG No	Emergency Response Guidebook - Number
Eye Dam.	Seriously damaging to the eye
Eye Irrit.	Irritant to the eye
Flam. Liq.	Flammable liquid
GHS	"Globally Harmonized System of Classification and Labelling of Chemicals" developed by the United Nations
IARC	International Agency for Research on Cancer
IATA	International Air Transport Association
IATA/DGR	Dangerous Goods Regulations (DGR) for the air transport (IATA)
ICAO	International Civil Aviation Organization
ICAO-TI	Technical instructions for the safe transport of dangerous goods by air
IMDG	International Maritime Dangerous Goods Code
IMDG-Code	International Maritime Dangerous Goods Code
LC50	Lethal Concentration 50%: the LC50 corresponds to the concentration of a tested substance causing 50 % lethality during a specified time interval
NPCA-HMIS® III	National Paint and Coatings Association: Hazardous Materials Identification System - HMIS® III, Third Edition

22. Safety Data Sheet

Safety Data Sheet

acc. to 29 CFR 1910.1200 App D

Kush Note

Version number: GHS 1.0 Date of compilation: 2024-05-26

Abbr.	Descriptions of used abbreviations
OSHA	Occupational Safety and Health Administration (United States)
PBT	Persistent, Bioaccumulative and Toxic
PNEC	Predicted No-Effect Concentration
ppm	Parts per million
Skin Corr.	Corrosive to skin
Skin Irrit.	Irritant to skin
Skin Sens.	Skin sensitization
STEL	Short-term exposure limit
STOT SE	Specific target organ toxicity - single exposure
TLV®	Threshold Limit Values
TWA	Time-weighted average
vPvB	Very Persistent and very Bioaccumulative

Key literature references and sources for data

OSHA Hazard Communication Standard (HCS), 29 CFR 1910.1200.

Transport of dangerous goods by road or rail (49 CFR US DOT). International Maritime Dangerous Goods Code (IMDG). Dangerous Goods Regulations (DGR) for the air transport (IATA).

Classification procedure

Physical and chemical properties: The classification is based on tested mixture.
Health hazards, Environmental hazards: The method for classification of the mixture is based on ingredients of the mixture (additivity formula).

List of relevant phrases (code and full text as stated in section 2 and 3)

Code	Text
H226	Flammable liquid and vapor.
H302	Harmful if swallowed.
H304	May be fatal if swallowed and enters airways.
H315	Causes skin irritation.
H317	May cause an allergic skin reaction.
H319	Causes serious eye irritation.
H335	May cause respiratory irritation.
H351	Suspected of causing cancer.

Disclaimer

This information is based upon the present state of our knowledge. This SDS has been compiled and is solely intended for this product.

23. Certificate of Analysis

SC Labs®

Hemp Quality Assurance Testing

CERTIFICATE OF ANALYSIS

DATE ISSUED 06/01/2025

SAMPLE DETAILS

SAMPLE NAME: Kush Cake - 7-508-9999
Other

CULTIVATOR / MANUFACTURER
Business Name:
License Number:
Address:

DISTRIBUTOR / TESTED FOR
Business Name: Eybna
License Number:
Address: 7647 Hayvenhurst Ave #30
Van Nuys CA 91406

SAMPLE DETAIL
Batch Number: BT25-0052
Sample ID: 250528L027

Date Collected: 05/28/2025
Date Received: 05/28/2025
Batch Size:
Sample Size: 1.0 unit
Unit Mass:
Serving Size:

Scan QR code to verify authenticity of results.

CANNABINOID ANALYSIS - SUMMARY

Total THC: Not Detected

Total CBD: Not Detected

Sum of Cannabinoids: Not Detected

Total Cannabinoids: Not Detected

Total THC/CBD is calculated using the following formulas to take into account the loss of a carboxyl group during the decarboxylation step:
Total THC = Δ^9-THC + (THCa (0.877))
Total CBD = CBD + (CBDa (0.877))
Sum of Cannabinoids = Δ^9-THC + THCa + CBD + CBDa + CBG + CBGa + THCV + THCVa + CBC + CBCa + CBDV + CBDVa + Δ^8-THC + CBL + CBN
Total Cannabinoids = (Δ^9-THC+0.877*THCa) + (CBD+0.877*CBDa) + (CBG+0.877*CBGa) + (THCV+0.877*THCVa) + (CBC+0.877*CBCa) + (CBDV+0.877*CBDVa) + Δ^8-THC + CBL + CBN

SAFETY ANALYSIS - SUMMARY

Pesticides: PASS **Residual Solvents:** PASS **Heavy Metals:** PASS

For quality assurance purposes. Not a Regulatory Hemp Lab Test Report. These results relate only to the sample included on this report. This report shall not be reproduced, except in full, without written approval of the laboratory.

Sample Certification: California Code of Regulations Title 4 Division 19. Department of Cannabis Control Business and Professions Code. Reference: Sections 26100, 26104 and 26110, Business and Professions Code.

Decision Rule: Statements of conformity (e.g. Pass/Fail) to specifications are made in this report without taking measurement uncertainty into account. Where statements of conformity are made in this report, the following decision rules are applied: PASS - Results within limits/specifications, FAIL - Results exceed limits/specifications.

References: limit of detection (LOD), limit of quantification (LOQ), not detected (ND), not tested (NT), µg/g = ppm, µg/kg = ppb

LQC verified by: Maria Garcia
Job Title: Senior Laboratory Analyst
Date: 06/01/2025

Approved by: Josh Wurzer
Job Title: Chief Compliance Officer
Date: 06/01/2025

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 CoA ID: 250528L027-001 Summary Page

23. Certificate of Analysis

SC Labs

Hemp Quality Assurance Testing

CERTIFICATE OF ANALYSIS

DATE ISSUED 06/01/2025

Cannabinoid Analysis

Tested by high-performance liquid chromatography with diode-array detection (HPLC-DAD).

Method: QSP 1157 - Analysis of Cannabinoids by HPLC-DAD

TOTAL THC: Not Detected

Total THC (Δ^9-THC+0.877*THCa)

TOTAL CBD: Not Detected

Total CBD (CBD+0.877*CBDa)

TOTAL CANNABINOIDS: Not Detected

Total Cannabinoids (Total THC) + (Total CBD) + (Total CBG) + (Total THCV) + (Total CBC) + (Total CBDV) + Δ^8-THC + CBL + CBN

TOTAL CBG: ND

Total CBG (CBG+0.877*CBGa)

TOTAL THCV: ND

Total THCV (THCV+0.877*THCVa)

TOTAL CBC: ND

Total CBC (CBC+0.877*CBCa)

TOTAL CBDV: ND

Total CBDV (CBDV+0.877*CBDVa)

CANNABINOID TEST RESULTS - 05/30/2025

COMPOUND	LOD/LOQ (mg/g)	MEASUREMENT UNCERTAINTY (mg/g)	RESULT (mg/g)	RESULT (%)
Δ^9-THC	0.002 / 0.014	N/A	ND	ND
Δ^8-THC	0.01 / 0.02	N/A	ND	ND
THCa	0.001 / 0.005	N/A	ND	ND
THCV	0.002 / 0.012	N/A	ND	ND
THCVa	0.002 / 0.019	N/A	ND	ND
CBD	0.004 / 0.011	N/A	ND	ND
CBDa	0.001 / 0.026	N/A	ND	ND
CBDV	0.002 / 0.012	N/A	ND	ND
CBDVa	0.001 / 0.018	N/A	ND	ND
CBG	0.002 / 0.006	N/A	ND	ND
CBGa	0.002 / 0.007	N/A	ND	ND
CBL	0.003 / 0.010	N/A	ND	ND
CBN	0.001 / 0.007	N/A	ND	ND
CBC	0.003 / 0.010	N/A	ND	ND
CBCa	0.001 / 0.015	N/A	ND	ND
SUM OF CANNABINOIDS			ND	ND

Pesticide Analysis

Pesticide and plant growth regulator analysis utilizing high-performance liquid chromatography-mass spectrometry (HPLC-MS) or gas chromatography-mass spectrometry (GC-MS).

*GC-MS utilized where indicated.

Method: QSP 1212 - Analysis of Pesticides and Mycotoxins by LC-MS or QSP 1213 - Analysis of Pesticides by GC-MS

PESTICIDE TEST RESULTS - 05/29/2025 PASS

COMPOUND	LOD/LOQ (µg/g)	ACTION LIMIT (µg/g)	MEASUREMENT UNCERTAINTY (µg/g)	RESULT (µg/g)	RESULT
Abamectin	0.03 / 0.10	0.3	N/A	ND	PASS
Azoxystrobin	0.02 / 0.07	40	N/A	ND	PASS
Bifenazate	0.01 / 0.04	5	N/A	ND	PASS
Bifenthrin	0.02 / 0.05	0.5	N/A	ND	PASS
Boscalid	0.03 / 0.09	10	N/A	ND	PASS
Chlorpyrifos	0.02 / 0.06	≥ LOD	N/A	ND	PASS
Cypermethrin	0.11 / 0.32	1	N/A	ND	PASS
Etoxazole	0.02 / 0.06	1.5	N/A	ND	PASS
Hexythiazox	0.02 / 0.07	2	N/A	ND	PASS
Imidacloprid	0.04 / 0.11	3	N/A	ND	PASS
Malathion	0.03 / 0.09	5	N/A	ND	PASS
Myclobutanil	0.03 / 0.09	9	N/A	ND	PASS
Permethrin	0.04 / 0.12	20	N/A	ND	PASS

Continued on next page

 CoA ID: 250528L027-001 Page 2 of 4

23. Certificate of Analysis

SC Labs

Hemp Quality Assurance Testing

CERTIFICATE OF ANALYSIS

DATE ISSUED 06/01/2025

Pesticide Analysis *Continued*

PESTICIDE TEST RESULTS - 05/29/2025 *continued* PASS

COMPOUND	LOD/LOQ (µg/g)	ACTION LIMIT (µg/g)	MEASUREMENT UNCERTAINTY (µg/g)	RESULT (µg/g)	RESULT
Piperonyl Butoxide	0.02 / 0.07	8	N/A	ND	PASS
Propiconazole	0.02 / 0.07	20	N/A	<LOQ	PASS
Spiromesifen	0.02 / 0.05	12	N/A	ND	PASS
Tebuconazole	0.02 / 0.07	2	N/A	<LOQ	PASS
Trifloxystrobin	0.03 / 0.08	30	N/A	ND	PASS

Residual Solvents Analysis

Residual Solvent analysis utilizing gas chromatography-mass spectrometry (GC-MS).

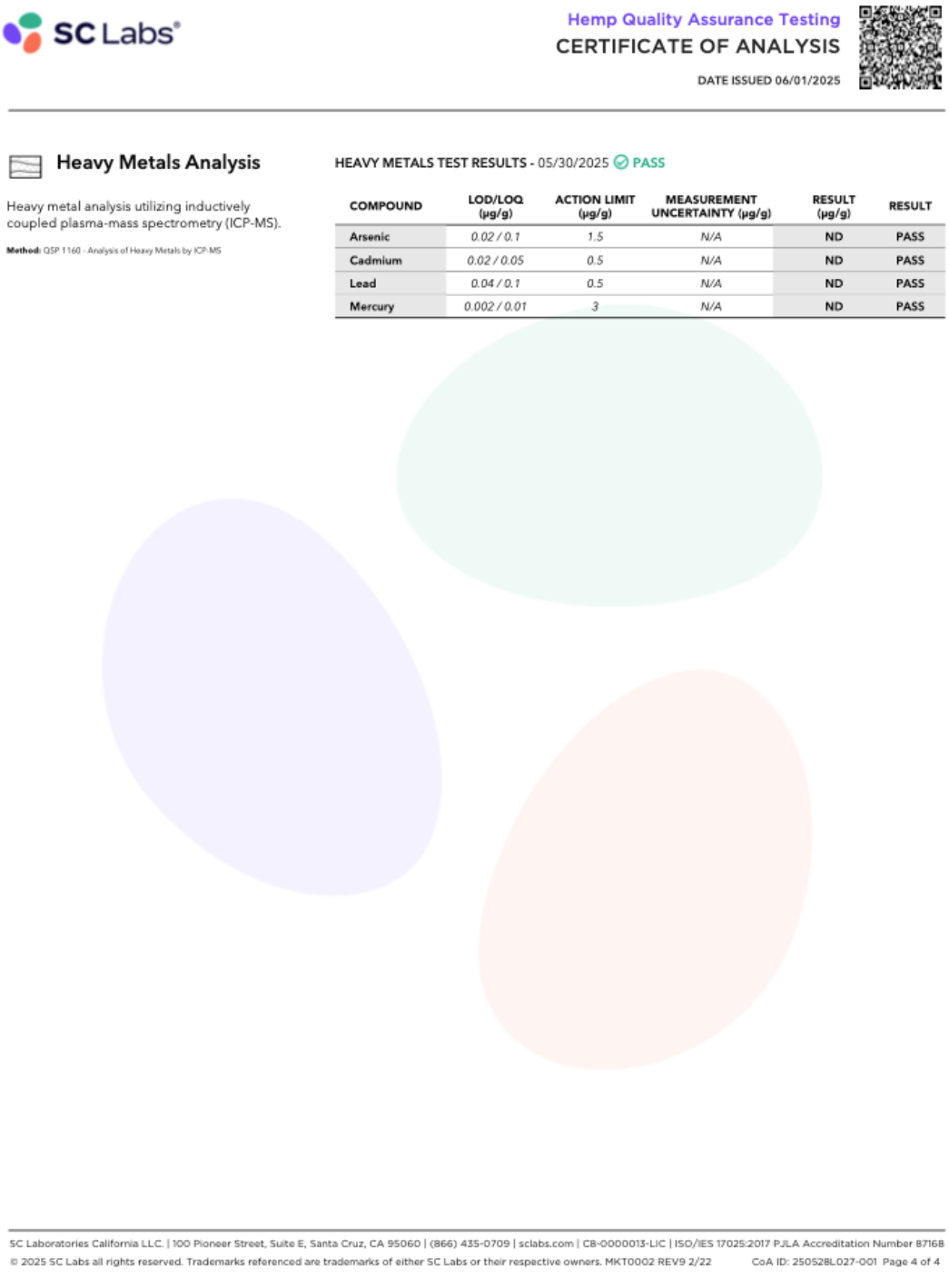
Method: QSP 1204 - Analysis of Residual Solvents by GC-MS

RESIDUAL SOLVENTS TEST RESULTS - 06/01/2025 PASS

COMPOUND	LOD/LOQ (µg/g)	ACTION LIMIT (µg/g)	MEASUREMENT UNCERTAINTY (µg/g)	RESULT (µg/g)	RESULT
Propane	10 / 20	5000	N/A	<LOQ	PASS
n-Butane	10 / 50	5000	N/A	ND	PASS
n-Pentane	20 / 50	5000	N/A	ND	PASS
n-Hexane	2 / 5	290	N/A	ND	PASS
n-Heptane	20 / 60	5000	N/A	ND	PASS
Benzene	0.03 / 0.09	1	N/A	ND	PASS
Toluene	7 / 21	890	N/A	ND	PASS
Total Xylenes	50 / 160	2170	N/A	ND	PASS
Methanol	50 / 200	3000	N/A	ND	PASS
Ethanol	20 / 50	5000	±2.5	85	PASS
2-Propanol (Isopropyl Alcohol)	10 / 40	5000	±1.8	66	PASS
Acetone	20 / 50	5000	±19.8	668	PASS
Ethyl Ether	20 / 50	5000	N/A	ND	PASS
Ethylene Oxide	0.3 / 0.8	1	N/A	ND	PASS
Ethyl Acetate	20 / 60	5000	N/A	ND	PASS
Chloroform	0.1 / 0.2	1	N/A	ND	PASS
Dichloromethane (Methylene Chloride)	0.3 / 0.9	1	N/A	ND	PASS
Trichloroethylene	0.1 / 0.3	1	N/A	ND	PASS
1,2-Dichloroethane	0.05 / 0.1	1	N/A	ND	PASS
Acetonitrile	2 / 7	410	N/A	ND	PASS

 CoA ID: 250528L027-001 Page 3 of 4

23. Certificate of Analysis



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Hemp Quality Assurance Testing

CERTIFICATE OF ANALYSIS

DATE ISSUED 06/01/2025

Heavy Metals Analysis

Heavy metal analysis utilizing inductively coupled plasma-mass spectrometry (ICP-MS).

Method: QSP 1160 - Analysis of Heavy Metals by ICP-MS

HEAVY METALS TEST RESULTS - 05/30/2025 PASS

COMPOUND	LOD/LOQ (µg/g)	ACTION LIMIT (µg/g)	MEASUREMENT UNCERTAINTY (µg/g)	RESULT (µg/g)	RESULT
Arsenic	0.02 / 0.1	1.5	N/A	ND	PASS
Cadmium	0.02 / 0.05	0.5	N/A	ND	PASS
Lead	0.04 / 0.1	0.5	N/A	ND	PASS
Mercury	0.002 / 0.01	3	N/A	ND	PASS

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CoA ID: 250528L027-001 Page 4 of 4

24. Label Example

25. Kosher Certificate

בס״ד

ORTHODOX UNION

LETTER OF KOSHER CERTIFICATION

UNION OF ORTHODOX JEWISH CONGREGATIONS OF AMERICA איחוד קהילות האורתודוקסים באמריקה
FORTY RECTOR STREET / NEW YORK, NY 10006 / 212-613-8241 / KOSHERLETTER@OU.ORG / OUKOSHER.ORG

June 04, 2025

This is to certify that the following product prepared by

Eybna Technologies, 1 HaYozma St., Kfar Saba, 4442214 ISRAEL

is under the supervision of the Kashruth Division of the Orthodox Union and is kosher as indicated below.

Product Name	UKD-ID	Status	Certification Requirements
Brand: Eybna			
• 10-556-9999 Red Raspberry	OUV3-5WY2SM7	Pareve	Ⓤ Symbol required.

Use of the OU trademark must comply with the terms set forth in a written agreement with the Orthodox Union. Any other use of the OU trademark is not authorized.

Rabbi Menachem Genack, *Rabbinic Adminstrator, CEO*

This certification is valid through 1/31/2026

Page 1 of 1

26. FSSC 22000 Certificate

Institute of Quality & Control

CERTIFICATE

IQC Institute of Quality & Control

NO. I29781

FSSC 22000

This is to certify that
The Food Safety Management System of

Eybna Technologies Ltd
1 Hayozma St. Kfar Saba, Israel

Has been assessed and determined to comply with the requirements of

FSSC 22000

Certification scheme for food safety management systems consisting of the following elements: ISO 22000:2018, ISO/TS 22002-1:2009 and Additional FSSC 22000 requirements (version 6).

This certificate is applicable for the scope of:
Mixing and packaging of taste and flavors formulations

Food Chain Subcategory K

Date of the last unannounced audit*:	NA
COID code	ISR-1-5243-682830
Certification decision date:	23.07.2024
Initial Certification date:	23.07.2024
Valid until:	22.07.2027

This certificate is subject to the continuing satisfactory operation of the Management System and periodic auditing by IQC

*At least one (1) surveillance audit is required to be undertaken unannounced after the initial certification audit and within each three (3) year period thereafter.

23.07.2024
Issue Date

Nir Halpern, CEO

IQC - Institute of Quality & Control
9 Shamira Imber Gadish st. Kiryat Ono, Israel
Tel: 03-9313555, Fax: 03-9044406
E-Mail: info@iqc.co.il, https://www.iqc.co.il

MGMT. SYS. RvA C 560

The authenticity of this certificate can be verified in the FSSC 22000 database of certified organizations available on www.fssc22000.com.

27. ISO 9001:2005 Certificate

Institute of Quality & Control

CERTIFICATE

IQC Institute of Quality & Control

No. I30297

This is to certify that
The Quality Management System of

Eybna Technologies Ltd
1 Hayozma St. Kfar Saba, Israel

Was audited by IQC and found to be
in compliance with the requirements of the standard:

ISO 9001:2015

This certificate is valid for the following scope of activities:

Research and development, production, marketing and sales of terpene-based extracts and products

This certificate is valid until:	08.12.2025
Certification cycle will end on:	08.12.2025
Date of previous certification cycle:	NA
Date of certification decision:	23.12.2024
Date of first approval:	22.09.2024

This certificate is subject to the continuing satisfactory operation
of the Management System and periodic auditing by IQC

26.12.2024
Issue Date

Nir Halpern, CEO

IQC - Institute of Quality & Control
9 Shamira Imber Gadish st. Kiryat Ono, Israel
Tel: 03-9313555
E-Mail: info@iqc.co.il, www.iqc.co.il

IAF MGMT. SYS. RvA C 560

28. GMP Certificate

www.health.gov.il

שרותי בריאות הציבור
שירות המזון הארצי
Public Health Service - Food Control Service

משרד הבריאות
לחיים בריאים יותר

To whom it may concern

Certificate of "Good Manufacturing Practice" (GMP)

Name of Food Business Operator(FBO):	**Eybna Technologies Ltd, 515146454**
Address:	**HaYozma St 1., Kfar Saba, Israel**
License number:	**103062**

This is to certify that in accordance with Article 42 on The Public Health Protection Food Law, 2015 (Israeli Food Law), and the regulation enacted under it, I hereby certify, based on the information available to me, that the FBO, meets our regulation of Good Manufacturing Practice (GMP). This certificate is subject to the requirements of the manufacturing license, including maintenance of appropriate sanitary and hygiene conditions during the manufacturing process and periodical inspections by the competent authority. In addition, have our permission to label food with the symbol of Good Manufacturing Practice, according to Article 46 of the Israeli Food Law.

The Certificate of Good Manufacturing Practice is granted for the following activities:
Manufacturing of flavor and fragrance formulations

Conditions of certificate:

1. This certificate is valid for this address of production only and it is not transferable.
2. The FBO must notify the District Food Control Services as early as possible and no later than 14 business days, of any changes to the application's details or to the enclosed documents regard this certificate.

This certificate is valid from: **24/02/2025** until: **24/02/2027**

בברכה,
פנינה אורן שניידור
ראשת שירות המזון הארצי
Pnina Oren-Shneidor
Head Food Control Service
Ministry Of Health, ISRAEL

Food Control Services
Ministry of Health
P.O.B 20301 Tel-Aviv 61203
Call.habriut@moh.health.gov.il
Tel *5400 **Fax**: 02-5655969

קול הבריאות
*5400

שירות המזון הארצי
משרד הבריאות
ת.ד. 20301 תל אביב 61203
Call.habriut@moh.health.gov.i
טל: *5400 פקס: 02-5655969