

Bulk SKU-1024 Low Temp Decarb THC Extract

Version 1.0.1 Last Modified: April 9, 2024

Product Name:

Minor RichTHC Extract SKU1024

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Product Information					
Product Label Name	High Purity Intermediary Minor Rich THC Extract - Rec Grade (SKU-1024)				
Brand(s)	GoodScienceRx				
Vendor	Protonify Corporati	on			
NNCP Number			Earliest Sa	ale Date	N/A
Class of Cannabis	Cannabis Extract		Intended Use Ir		Inhalation
Cannabis Product Form	Extract / Concentrate		Discrete Units N		N/A
Accessory Description	N/A			•	
Net Weight/Volume per unit	N/A Ur		Units per	Container	N/A
Net Weight/Volume per Container (accepted range)	1,000 mg +/- 50 mg	9			
Cannabinoid Content	Tetrahydrocannabinol (THC)		Cannabidiol (CBD)		
Garriabiliola Goriterit	Excluding THCA	Total THC		Excluding CBDA	
Per Unit	N/A	N/A		N/A	N/A
Cannabinoid Summary (mg/g) (accepted range)	730 mg/g to 900 mg/g	825 mg/g +/-10% (765 mg/g to 935 mg/g)		50 mg/g	50 mg/g +/-100% (0 mg/g to 100 mg/g)
Other Cannabinoids	CBG, CBC, CDB, THCv				
	Texture	Colour		Consistency	Other
Physical Attributes	Liquid with high	yellow to reddish		consistent	N/A
	viscosity	brown		color, potency	,
				viscosity	
_	Scent/Aroma Faint Cannabis Smell				
Sensory Attributes					



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Packaging Description				
Reference Image	Primary Packaging Total Grosson To			
Description	Stainless Steel Vault or FDA Compliant Amber Glass Bottle			
Technical Specifications	Cap: Black Phenol (FDA compliant) Temperature: Application: -51° to 82° C (-60° to 180° F) Usage: Leakproof and airtight.			
Dimensions	Container: Amber Glass, 16 Oz, 32 Oz. Vault, 3L, 6L, 12L			
Supplier	Uline			
Manufacturer	Uline			
Mandatory Label requirement	"PRODUCT TESTED" Unique Container ID Batch # Cultivar/Product Name Cannabis Type Packaging Date Net weight Gross weight Tare weight Supervisor (second set of eyes) initials			
Mandatory bulk	FDA grade container			
packaging requirement	-			
Additional requirements API Information	Store in a cool dry place out of direct sunlight.			

API Information	on
Ingredient Declaration	THC
Carrier Type	None
Terpenes / Flavouring	Negligible, as the majority of Terpenes are removed during the manufacturing process and not re-captured for processing.
Restricted Ingredients	None
Extraction / Process Method	Liquid Liquid Extraction, chromatography based purification and vacuum based crystallization.
Species / Strain Requirements	Strain agnostic
Input Cannabis Potency (acceptable range)	220 mg/g; based on oil density 0.220 g/mL (180 mg/g to 240 mg/g)



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Vendor Protonify approved supplier

Recipe / Ingredient Information						
Ingredient Name	Manufacturer (inc. facility, if applicable)	Manufacturer Product Name	% / wt. in recipe	Notes		
Pure Intermediary Minor Rich THC Extract	Protonify	SKU-1024	100%	N/A		

Other Information	
Processing Aid(s)	Product Manufacturer: Manufacturer Product Name: Isopropyl Manufacturer: Manufacturer: Heptane
Additives	N/A
Storage Condition	Cool,dry area out of direct sunlight

Allergen Information					
Label Claim	Co	ntains:	N/A		
Laber Claim	May Contain: NA				
			t in Product(s)		
Allergen	Present in Product		ufactured on ame Line	Present in Facility	Ingredient Carryover
Eggs	N		Ζ	N	NA
Milk	N		Ν	N	NA
Mustard	N		N	N	NA
Peanuts	N		N	N	NA
Crustaceans and molluscs	N		Ν	N	NA
Fish	N		N	N	NA
Sesame seeds	N		N	N	NA
Soy	N		N	N	NA
Sulphites	N		N	N	NA
Tree nuts	N		N	N	NA



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Shipping Case Information				
Shipping Container / Case Material	Secondary Containment FDA grade plastic bag Uline Cardboard boxes with packing material. Box within a box, with extra spaces filled with dummy / empty boxes if necessary. Sealed with tamper proof tape Contains Shipping Collateral Package: Manifest (container Unique ID, Tare, Gross, Net material weight), box unique ID with total count in shipment, PO reference, QA contact / approval), Certificate of Manufacture, Certificate of Analysis, Chain of Custody (signable and return form), MSDS & Source Biomass pesticide checklist			
Dimension	Dimensions- 22x22x22 but can be packaged in larger or smaller qty by customer request.			
Minimum Label Requirements	Toteof Batch # and Product Name Cannabis Type Packaging Date Net weight Gross weight			
Tamper evidence	Individual containers sealed and boxes sealed with tamper proof tape.			



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Analytical / Laboratory Release Specif		Mathad/Dafarrana **
Test Description	Limit Report as is (mg/g)	Method/Reference**
THC	. , , , , , , , , , , , , , , , , , , ,	4
THCA	Report as is (mg/g)	4
Total THC	See Product Information Section	
CBD	Report as is (mg/g)	HPLC-DAD:
CBDA	Report as is (mg/g)	Perkin Elmer LC300 ORG-M-071
Total CBD	See Product Information Section	ONG-WI-07 I
CBG	Report as is (mg/g)	1
CBGA	Report as is (mg/g)	7
	1 (30)	
Pesticides (tested on flower / dry input)	Absent* (limits defined by Health Canada)	LC-MS/MS Dual Ion Source. Limits are set by Health Canada for Cannabis concentrate ORG-M-052
Aflatoxin B1	< 2ppb	LC-MS/MS ESI
Total Aflatoxin (B _{1.} B _{2.} G _{1.} G ₂)	< 4ppb	EP 2.8.22 EP 2.8.18/USP<561> ORG-M-069
Total Aerobic Bacterial Count (TAMC)	< 200 cfu/g or cfu/mL (Inhalation) < 50,000 cfu/g or cfu/mL (Ingestion)	
Total Yeast & Mold (TYMC)	< 20 cfu/g or cfu/mL (Inhalation) < 500 cfu/g or cfu/mL (Ingestion)	EP 2.6.12/USP<61> EP 2.6.13/USP<62>
Bile Tolerant Gram-Negative Bacteria (BTGN)	Absent in 1g or 1ml (Inhalation) < 100 cfu/g or cfu/mL (Ingestion)	EP 2.6.31/USP<62>
Escherichia coli	Absent in 1g or 1ml (Ingestion)	EP 5.1.4 Table 1 (Inhalation)
Salmonella spp.	Absent in 25g or 25ml (Ingestion)	EP 5.1.8 Table B (Ingestion)
S. aureus	Absent in 1g or 1ml (Inhalation)	
P. aeruginosa.	Absent in 1g or 1ml (Inhalation)	7
Ochratoxin A	< 20 ppb	LC-MS/MS ESI EP 2.8.22 EP 2.4.20/USP<233>
Residual Solvent: Ethanol Isopropanol Heptane	< 5,000ppm for class 2 solvents	GC-FID USP <467> ORG-M-074 ICH Topic Q3C (R6) (R8)–Table 3
Methanol	< 3000 ppm for Class 3	GC-FID USP <467> ORG-M-074 ICH Topic Q3C (R6) (R8)–Table 3
Arsenic (As)	< 0.2ppm (Inhalation)	IPC-MS
Cadmium (Cd)	< 0.3ppm (Inhalation)	USP <232> Table 3
Lead (Pb)	< 0.5ppm (Inhalation)	EP 5.20 ICHQ3
` '	< 0.1ppm (Inhalation)	USP<233>/E.P 2.4.20

^{**} **Methods** and **Validation Summaries** are to be provided during an Audit (see Quality Agreement). Any changes to methods, validations and labs necessitates notification per the Change Control requirements