### **High Performance Liquid Analytics**

111 Westport Plaza Dr. 6th Floor St. Louis, MO. 63146

## **Certificate** of Analysis

June 15, 2020 Derma Thereal **SAFETY RESULTS**  For informational purposes only.

Product Name: Eye Cream 20MG Manufacturer: DermaThereal

Type:Topical

Test: Non-Compliance Full-Panel

Sample ID:-T079 Sample Size: 5ml

Retail Product Size:20mg/15ml/.5oz

Ordered: 06/06/20 Sampled: 06/07/20 Completed: 06/15/20

Expires: 06/15/21

Sampling Method: SOP Client Method



PRODUCT IMAGE

PASSED

Page 1 of 5

MISC.

Pesticides Heavy Metals

Microbials

Mycotoxins

Residuals

Filth

Water Activity

Moisture

Terpenes

PASSED PASSED

PASSED

PASSED

PASSED

PASSED NOT TESTED NOT TESTED

TESTED

#### CANNABINOID RESULTS

**Total THC** 0.0% **Total CBD** 

.75mg/ml

CBD/Container: 22.40mg

**Total Cannabinoids** 

22.40ma

Total Cannabinoids/Container: 22.40mg

CBC	CBGA	CBG	CBN	THCV	D-8 THC	CBDV	CBDA	CBD	D-9 THC	THCA
ND	ND	ND	ND	ND	ND	ND	0.05	3.73	ND	ND
mg/v	mg/v	mg/v	mg/v	mg/v	mg/v	mg/v	mg/v	mg/v	mg/v	mg/v
ND	ND	ND	ND	ND	ND	ND	0.01	0.75	ND	ND
mg/v	mg/v	mg/v	mg/v	mg/v	mg/v	mg/v	mg/v	mg/v	mg/v	mg/v
0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01
LOD	LOD	LOD	LOD	LOD	LOD	LOD	LOD	LOD	LOD	LOD

Foreign Matter PASSED

Analyzed By Weight Date ID-710

25ml 06/12/20

LOD(ppm) N/A

Reviewed On - 06/15/20

Instrument Used: Filth/Foreign Material Microscope

This includes but is not limited to hair, insects, feces, packaging contaminants, and manufacturing waste and by-products.

#### **Cannabinoid Profile Test**

Analyzed By ID-710 5ml

Weight

**Extraction date** 

06/07/20

**Extracted By** 

ID-710

Reviewed On - 06/15/20 Instrument Used: HPLA-1100-DAD-#01

Reagent 030520.03

Dilution 400

Full spectrum cannabinoid analysis utilizing High Performance Liquid Chromatography with UV detection (HPLC-UV).

This report shall not be reproduced, unless in its entirety, without written approval from High Performance Liquid Analytics. This report is a High Performance Liquid Analytics certification. The results relate only to the material or product analyzed. Test results are confidential unless explicitly waived otherwise. Void after 1 year from test end date Cannabinoid content of batch material may vary depending on sampling error. IC=In-control QC parameter, NC=Non-controlled QC parameter, ND=Not Detected, NA=Not Analyzed, ppm=Parts Per Million, ppb=Parts Per Billion. Limit of Detection (LoD) and Limit Of Quantitation (LoQ) are terms used to describe the smallest concentration that can be reliably measured by an analytical procedure. RPD=Repro-ducibility of two measurements. Action Levels are State determined thresholds for human safety for consumption and/or inhalation. The result >99% are variable based on uncertainty of measurement (UM) for the analyte. The UM error is available from the lab upon request. The "Decision Rule" for the pass/fail does not include the UM. The limits are based on F.S. Rule 64-4.310.

C. Kukowski LAB DIRECTOR



### **High Performance Liquid Analytics**

111 Westport Plaza Dr. 6th Floor St. Louis, MO. 63146

**Certificate of Analysis** 

Derma Thereal

Product Name: Eye Cream 20MG Manufacturer: DermaThereal

Type:Topical

Test: Non-Compliance Full-Panel

Sample ID:-T079 Sample Size: 5ml

Retail Product Size:20mg/30ml/1oz

Ordered: 06/06/20 Sampled: 06/07/20

Completed: 06/15/20 Expires: 06/15/21

Sampling Method: SOP Client Method

**D5 GG9 8** 

Page 2 of 5

### **Terpenes**

HI	gnyx

HYfdYbYg"	@C8	Ib]hg"	FYgi `hfl I
5 @D<5!798F9B9C	0.007	%	ND
5 @D<5! <i @b9<="" ai="" th=""><th>0.007</th><th>%</th><th>ND</th></i>	0.007	%	ND
5 @D<5!D=B9B9'	0.007	%	ND
5 @D<5!H9FD=B9B9	0.007	%	ND
69H5!AMF79B9	0.007	%	ND
69H5!D=B9B9	0.007	%	ND
6CFB9C@	0.013	%	ND
75AD<9B9	0.007	%	ND
75AD <cf< th=""><th>0.013</th><th>%</th><th>ND</th></cf<>	0.013	%	ND
75FMCD <m@@b9'cl-89'< th=""><th></th><th></th><th></th></m@@b9'cl-89'<>			
798FC@	0.007	%	ND
5 @D<5!6=G56C@C@	0.007	%	ND
G5 6 ±B9 B9	0.007	%	ND
G5 6 =B9 B9 '< M8 F5 H9	0.007	%	ND
H9 F D=B9 C @	0.007	%	ND
H9 F D=BC @ B9	0.007	%	ND
69H5!75FMCD <m@@b9< th=""><th>0.007</th><th>%</th><th>ND</th></m@@b9<>	0.007	%	ND
HF5 BG!B9 FC @8 C @	0.007	%	ND
J5 @ B7 9 B9 '	0.007	%	ND
DI @9; CB9	0.007	%	ND
5 @D<5!D<9 @@5 B8 F9 B9	0.007	%	ND
C7=A9B9	0.007	%	ND
B9FC@	0.007	%	ND
@B5 @CC @	0.007	%	ND
@ACB9B9	0.007	%	ND
; I 5=C@	0.007	%	ND
; 9F5BM@579H5H9	0.007	%	ND
; 9F5B€C@	0.007	%	ND
; 5AA5!H9FD=B9B9	0.007	%	ND
: 9B7 < CB9	0.007	%	ND
: 5 FB9 G9 B9	0.007	%	ND
TPRI.		\$.	
HcHU.		Þ	

HYfdYbYg <sup>*</sup>	@C8	I b]ltg"	FYgi `hfl Ł
91 75 @MDHC@	0.007	%	ND
=GC6 CFB9 C@	0.007	%	ND
<9L5 <m8fch<mac@< th=""><th>0.007</th><th>%</th><th>ND</th></m8fch<mac@<>	0.007	%	ND
: 9B7 < M@5 @7C < C@	0.007	%	ND
'!75F9B9	0.007	%	ND
7 =G!B9 F C @8 C @	0.007	%	ND
=GCDI @9; C@	0.007	%	ND

#### **Terpenes**

H9 GH9 8

5 bUmnYX'Vm KY][\h 91 HUWich XUNY '91 HUWYX'6 m

ID-710

1.0045g

06/08/20

ID-710

FYj ]Yk YX'Cb - 06/15/20

⇒bglfi a Ybhil gYX'. Liquid Injection GCMS QP2010

FYU[Ybhi	8] i l]cb	7 cbgi a g"≇8 <sup>·</sup>
021420.11	10	180111
012120.R13		280653964

Terpenoid profile screening is performed using GC-MS with Liquid Injection (Gas Chromatography - Mass Spectrometer) which can screen 38 terpenes using Method SOP.T.40.091 Terpenoid Analysis Via GC/MS.

This report shall not be reproduced, unless in its entirety, without written approval from High Performance Liquid Analytics. This report is a High Performance Liquid Analytics certification. The results relate only to the material or product analyzed. Test results are confidential unless explicitly waived otherwise. Void after 1 year from test end date. Cannabinoid content of batch material may vary depending on sampling error. IC=In-control QC parameter, NC=Non-controlled QC parameter, ND=Not Detected, NA=Not Analyzed, ppm=Parts Per Million, ppb=Parts Per Billion. Limit of Detection (LoD) and Limit Of Quantitation (LoQ) are terms used to describe the smallest concentration that can be reliably measured by an analytical procedure. RPD=Repro-ducibility of two measurements. Action Levels are State determined thresholds for human safety for consumption and/or inhalation. The result >99% are variable based on uncertainty of measurement (UM) for the analyte. The UM error is available from the lab upon request. The "Decision Rule" for the pass/fail does not include the UM. The limits are based on F.S. Rule 64-4.310.

C. Kukowski LAB DIRECTOR



# High Performance Liquid Analytics

111 Westport Plaza Dr. 6th Floor St. Louis, MO. 63146

## **Certificate of Analysis**

Derma Thereal

Product Name: Eye Cream 20MG Manufacturer: DermaThereal

Type:Topical

Test: Non-Compliance Full-Panel

Sample ID:-T079 Sample Size: 5ml

Retail Product Size:20mg/30ml/1oz

Ordered: 06/06/20 Sampled: 06/07/20 Completed: 06/15/20

Expires: 06/15/21

Sampling Method: SOP Client Method

**PASSED** 

Page 3 of 5

### **Pesticides**

#### **PASSED**

Pesticides	LOD	Units	Action Level	Res		
ABAMECTIN B1A	0.01	ppm	0.3	ND		
ACEPHATE	0.01	ppm	3	ND		
ACEQUINOCYL	0.01	ppm	2	ND		
ACETAMIPRID	0.01	ppm	3	ND		
ALDICARB	0.01	ppm	0.1	ND		
AZOXYSTROBIN	0.01	ppm	3	ND		
BIFENAZATE	0.01	ppm	3	ND		
BIFENTHRIN	0.01	ppm	0.5	ND		
BOSCALID	0.01	ppm	3	ND		
CAPTAN	0.07	ppm	3	ND		
CARBARYL	0.05	ppm	0.5	ND		
CARBOFURAN	0.01	ppm	0.1	ND		
CHLORANTRANILIPROLE	0.1	ppm	3	ND		
CHLORFENAPYR	0.01	ppm	0.1	ND		
CHLORMEQUAT CHLORIDE	0.05	ppm	3	ND		
CHLORPYRIFOS	0.01	ppm	0.1	ND		
CLOFENTEZINE	0.02	ppm	0.5	ND		
COUMAPHOS	0.01	ppm	0.1	ND		
CYFLUTHRIN	0.05	ppm	1	ND		
CYPERMETHRIN	0.05	ppm	1	ND		
DAMINOZIDE	0.01	ppm	0.1	ND		
DIAZANON	0.01	ppm	0.2	ND		
DICHLORVOS	0.01	ppm	0.1	ND		
DIMETHOATE	0.01	ppm	0.1	ND		
DIMETHOMORPH	0.02	ppm	3	ND		
ETHOPROPHOS	0.01	ppm	0.1	ND		
ETOFENPROX	0.01	ppm	0.1	ND		
ETOXAZOLE	0.01	ppm	1.5	ND		
FENHEXAMID	0.01	ppm	3	ND		
FENOXYCARB	0.01	ppm	0.1	ND		
FENPYROXIMATE	0.01	ppm	2	ND		
FIPRONIL	0.01	ppm	0.1	ND		
FLONICAMID	0.01	ppm	2	ND		
FLUDIOXONIL	0.01	ppm	3	ND		
HEXYTHIAZOX	0.01	ppm	2	ND		
IMAZALIL	0.01	ppm	0.1	ND		
IMIDACLOPRID	0.04	ppm	3	ND		
KRESOXIM-METHYL	0.01	ppm	1	ND		
MALATHION	0.02	ppm	2	ND		
METALAXYL	0.01	ppm	3	ND		
METHIOCARB	0.01	ppm	0.1	ND		

t	Pesticides	LOD	Units	Action Level	Result
	METHOMYL	0.01	ppm	0.1	ND
	METHYL PARATHION	0.005	ppm	0.1	ND
	MEVINPHOS	0.01	ppm	0.1	ND
	MYCLOBUTANIL	0.01	ppm	3	ND
	NALED	0.025	ppm	0.5	ND
	OXAMYL	0.05	ppm	0.5	ND
	PACLOBUTRAZOL	0.01	ppm	0.1	ND
	PHOSMET	0.01	ppm	0.2	ND
	PIPERONYL BUTOXIDE	0.1	ppm	3	ND
	PRALLETHRIN	0.01	ppm	0.4	ND
	PROPICONAZOLE	0.01	ppm	1	ND
	PROPOXUR	0.01	ppm	0.1	ND
	PYRETHRINS	0.05	ppm	1	ND
	PYRIDABEN	0.02	ppm	3	ND
	SPINETORAM	0.02	ppm	3	ND
	SPIROMESIFEN	0.01	ppm	3	ND
	SPIROTETRAMAT	0.01	ppm	3	ND
	SPIROXAMINE	0.01	ppm	0.1	ND
	TEBUCONAZOLE	0.01	ppm	1	ND
	THIACLOPRID	0.01	ppm	0.1	ND
	THIAMETHOXAM	0.05	ppm	1	ND
	TOTAL CONTAMINANT LOAD	0	ppm	20	ND
	(PESTICIDES)				
	TOTAL PERMETHRIN	0.01	ppm	1	ND
	TOTAL SPINOSAD	0.01	ppm	3	ND

#### **Pesticides**

#### **PASSED**

Analyzed by	Weight	Extraction date	Extracted By
ID-710	1.0501g	06/08/20	ID-710

0.01

Reviewed On - 06/15/20

**TRIFLOXYSTROBIN** 

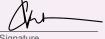
sult

Instrument Used: Liquid Injection GCMS QP2010

Reagent	Dilution	Consums. ID
013120.28	10	180111
031220.R10		280653964
032320.R17		

This report shall not be reproduced, unless in its entirety, without written approval from High Performance Liquid Analytics. This report is a High Performance Liquid Analytics certification. The results relate only to the material or product analyzed. Test results are confidential unless explicitly waived otherwise. Void after 1 year from test end date. Cannabinoid content of batch material may vary depending on sampling error. IC=In-control QC parameter, NC=Non-controlled QC parameter, ND=Not Detected, NA=Not Analyzed, ppm=Parts Per Million, pb=Parts Per Billion. Limit of Detection (LoD) and Limit Of Quantitation (LoQ) are terms used to describe the smallest concentration that can be reliably measured by an analytical procedure. RPD=Repro-ducibility of two measurements. Action Levels are State determined thresholds for human safety for consumption and/or inhalation. The result >99% are variable based on uncertainty of measurement (UM) for the analyte. The UM error is available from the lab upon request. The "Decision Rule" for the pass/fail does not include the UM. The limits are based on F.S. Rule 64-4.310.

C. Kukowski LAB DIRECTOR



06-15-2020

Signed On

## High Performance Liquid Analytics

111 Westport Plaza Dr. 6th Floor St. Louis, MO. 63146

**Certificate of Analysis** 

Derma Thereal

Product Name: Eye Cream 20MG
Manufacturer: DermaThereal
Type:Topical
Test: Non-Compliance Full-Panel

Sample ID:-T079 Sample Size: 5ml

Retail Product Size:20mg/15ml/.5oz Ordered: 06/06/20

Sampled: 06/07/20 Completed: 06/15/20

Expires: 06/15/21 Sampling Method: SOP Client Method

**PASSED** 

Page 4 of 5

#### **Residual solvents**

**PASSED** 

Solvent	LOD	Units	Action Level	Pass/Fa	il Result
			(PPM)		
1,1-DICHLOROETHENE	1	ppm	8	PASS	ND
1,2-DICHLOROETHANE	0.18	ppm	2	PASS	ND
2-PROPANOL	45	ppm	500	PASS	ND
ACETONE	67.5	ppm	750	PASS	ND
ACETONITRILE	5.4	ppm	60	PASS	ND
BENZENE	0.09	ppm	1	PASS	ND
BUTANES (N-BUTANE)	96	ppm	5000	PASS	ND
CHLOROFORM	0.18	ppm	2	PASS	ND
DICHLOROMETHANE	3.75	ppm	125	PASS	ND
ETHANOL	90	ppm	000000	PASS	776.749
ETHYL ACETATE	36	ppm	400	PASS	ND
ETHYL ETHER	45	ppm	500	PASS	ND
ETHYLENE OXIDE	0.6	ppm	5	PASS	ND
HEPTANE	45	ppm	5000	PASS	ND
METHANOL	22.5	ppm	250	PASS	ND
N-HEXANE	4.5	ppm	250	PASS	ND
PENTANES (N-PENTANE)	67.5	ppm	750	PASS	ND
PROPANE	120	ppm	5000	PASS	ND
TOLUENE	13.5	ppm	150	PASS	ND
TOTAL XYLENES	13.5	ppm	150	PASS	ND
TRICHLOROETHYLENE	2.25	ppm	25	PASS	ND

Analyzed by Weight Extraction date Extracted By
1D-710 0.0294g 06/08/20 1D-710

**Reviewed On** - 06/15/20

Instrument Used: Headspace GCMS

Reagent Dilution Consums. ID

1 00279984
161291-1
24154107

This report shall not be reproduced, unless in its entirety, without written approval from High Performance Liquid Analytics. This report is a High Performance Liquid Analytics certification. The results relate only to the material or product analyzed. Test results are confidential unless explicitly waived otherwise. Void after 1 year from test end date. Cannabinoid content of batch material may vary depending on sampling error. IC=In-control QC parameter, NC=Non-controlled QC parameter, ND=Not Detected, NA=Not Analyzed, ppm=Parts Per Million, ppb=Parts Per Billion. Limit of Detection (LoD) and Limit Of Quantitation (LoQ) are terms used to describe the smallest concentration that can be reliably measured by an analytical procedure. RPD=Repro-ducibility of two measurements. Action Levels are State determined thresholds for human safety for consumption and/or inhalation. The result >99% are variable based on uncertainty of measurement (UM) for the analyte. The UM error is available from the lab upon request. The "Decision Rule" for the pass/fail does not include the UM. The limits are based on F.S. Rule 64-4.310.

C. Kukowski LAB DIRECTOR



06-15-2020

Signed On

# High Performance Liquid Analytics

111 Westport Plaza Dr. 6th Floor St. Louis, MO. 63146

## **Certificate of Analysis**

Derma Thereal

Product Name: Eye Cream 20MG Manufacturer: DermaThereal

Type:Topical

Test: Non-Compliance Full-Panel

Sample ID:-T079
Sample Size: 5ml

Retail Product Size:20mg/15ml/.5oz

Ordered: 06/06/20 Sampled: 06/07/20 Completed: 06/15/20

Expires: 06/15/21
Sampling Method: SOP Client Method

**PASSED** 

Page 5 of 5

### **Mycotoxins**

#### **PASSED**

Analyte	LOD	Units	Result	Action Level (PPM)
AFLATOXIN G2	0.002	ppm	ND	0.02
AFLATOXIN G1	0.002	ppm	ND	0.02
<b>AFLATOXIN B2</b>	0.002	ppm	ND	0.02
AFLATOXIN B1	0.002	ppm	ND	0.02
OCHRATOXIN A+	0.002	ppm	ND	0.02

Aflatoxins B1, B2, G1, G2, and Ochratoxins A testing using LC-MS. (Method: SOP.T.30.065 for Sample Preparation and SOP.T40.065 Procedure for Mycotoxins Quantification Using LCMS. LOQ 1.0 ppb). Aflatoxin B1, B2, G1, and G2 must individually be <20ug/Kg. Ochratoxins must be <20ug/Kg.

#### **Heavy Metals**

#### **PASSED**

**Dilution** 

**Extracted By** 

ID-710

Reagent	
032420.R01	
031820.R03	
031820.R02	
031920.R01	
111319.02	

Analyzed by

Reviewed On - 06/15/20 Instrument Used: ICPMS-2030 B

ID-710

Metal LOD Unit Result Action Level (PPM)

**Extraction date** 

06/07/20

ARSENIC	0.02	ppm	ND	.5
CADMIUM	0.02	ppm	ND	0.5
LEAD	0.05	ppm	ND	0.5
MERCURY	0.02	ppm	ND	3

Weight

0.2808g

#### **Microbials**

#### **PASSED**

nalyte	Res
SPERGILLUS_FLAVUS	not present in 1 gra

ASPERGILLUS\_FUMIGATUS not present in 1 gram.
ASPERGILLUS\_NIGER not present in 1 gram.
ASPERGILLUS\_TERREUS not present in 1 gram.
ESCHERICHIA\_COLI\_SHIGELLA\_SPP not present in 1 gram.
SALMONELLA\_SPECIFIC\_GENE not present in 1 gram.

SALMONELLA\_SPECIFIC\_GENE not present in 1 gram.

TOTAL\_YEAST\_AND\_MOLD <100

**Reviewed On** - 06/15/20

Instrument Used: PathogenDX PCR\_Array Scanner, PathogenDX PCR\_DA-171

Analyzed by Weight Extraction date Extracted By

ID-710 1.0197g 06/07/20 ID-710

Reagent Dilution Consums. ID
181019-274

Microbiological testing for Fungal and Bacterial Identification via Polymerase Chain Reaction (PCR) method consisting of sample DNA amplified via tandem Polymerase Chain Reaction (PCR) as a crude lysate which avoids purification. (Method SOP.T.40.043) If a pathogenic Escherichia Coli, Salmonella, Aspergillus fumigatus, Aspergillus flavus, Aspergillus niger, or Aspergillus terreus is detected in 1g of a sample, the sample fails the microbiological-impurity testing.

This report shall not be reproduced, unless in its entirety, without written approval from High Performance Liquid Analytics. This report is a High Performance Liquid Analytics certification. The results relate only to the material or product analyzed. Test results are confidential unless explicitly waived otherwise. Void after 1 year from test end date. Cannabinoid content of batch material may vary depending on sampling error. IC=In-control QC parameter, NC=Non-controlled QC parameter, ND=Not Detected, NA=Not Analyzed, ppm=Parts Per Million, ppb=Parts Per Billion. Limit of Detection (LoD) and Limit Of Quantitation (LoQ) are terms used to describe the smallest concentration that can be reliably measured by an analytical procedure. RPD=Repro-ducibility of two measurements. Action Levels are State determined thresholds for human safety for consumption and/or inhalation. The result >99% are variable based on uncertainty of measurement (UM) for the analyte. The UM error is available from the lab upon request. The "Decision Rule" for the pass/fail does not include the UM. The limits are based on F.S. Rule 64-4.310.

C. Kukowski

