



ISANDO 20

Certificate of Analysis

Certificate of Analysis

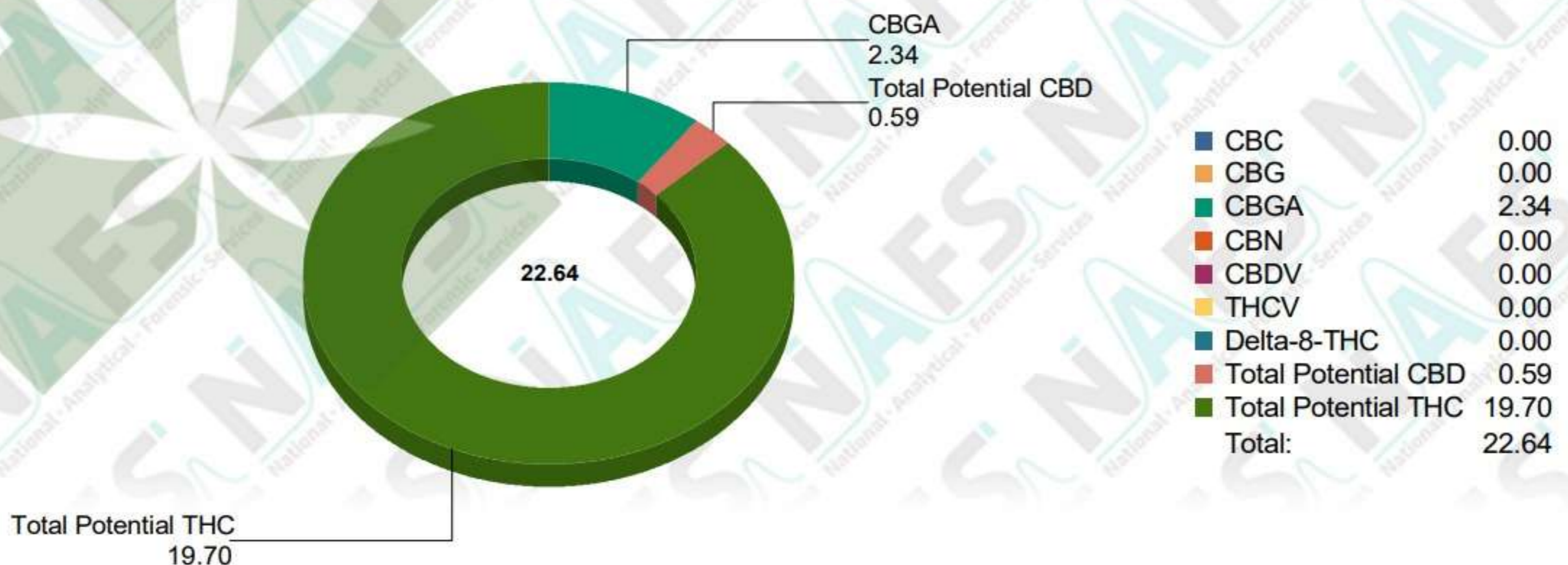
Sample ID: 220831-004-001
Retention Sample ID: 11,973
Client (Code) Name: (CAN0120) MedCan (Pty) Ltd
Client Address: S10 Euro Centre, 363 Rivonia Boulevard, Rivonia
Batch: NA-0005-220822-1
Date Receipt: 9/1/2022 9:45:39AM
Date Report Released: 9/13/2022 3:18:32PM
Sample Type: Plant Material
Package Condition: Intact
Received By: JLEYGONIE
Sample Amount: 3 gram
Tamper Proof: No



Test Name: TM006.3 (Potency Cannabis HPLC) (220831-004-001)
Specification: N/A
Test Method: *TM006.3
Sampling Method: SH007

Analyte	Results	*Std Dev.	*Spec Limits
CBC	ND % Weight	% Weight	N/A
CBD	0.595 % Weight	+/- 0.019 % Weight	N/A
CBD-A	ND % Weight	% Weight	N/A
CBD-V	ND % Weight	% Weight	N/A
CBG	ND % Weight	% Weight	N/A
CBG-A	2.341 % Weight	+/- 0.240 % Weight	N/A
CBN	ND % Weight	% Weight	N/A
Delta-8-THC	ND % Weight	% Weight	N/A
Delta-9-THC	0.616 % Weight	+/- 0.048 % Weight	N/A
THC-A	21.765 % Weight	+/- 2.266 % Weight	N/A
THC-V	ND % Weight	% Weight	N/A
Total Potential CBD	0.595 % Weight	+/- 0.019 % Weight	N/A
Total Potential THC	19.703 % Weight	+/- 2.036 % Weight	N/A

Analyst: LMARAIS 2022/09/09 10:34



*ND (Not Detected)
*The current test method version is Validated
*Standard Deviation (Uncertainty of measurement of applicable duplicate sample)
*Specification Limits (Specified according to pharmacopeial or client specification limits where applicable)

Medcon



Test Name: TM003.5 (Terpenes) (220831-004-001)

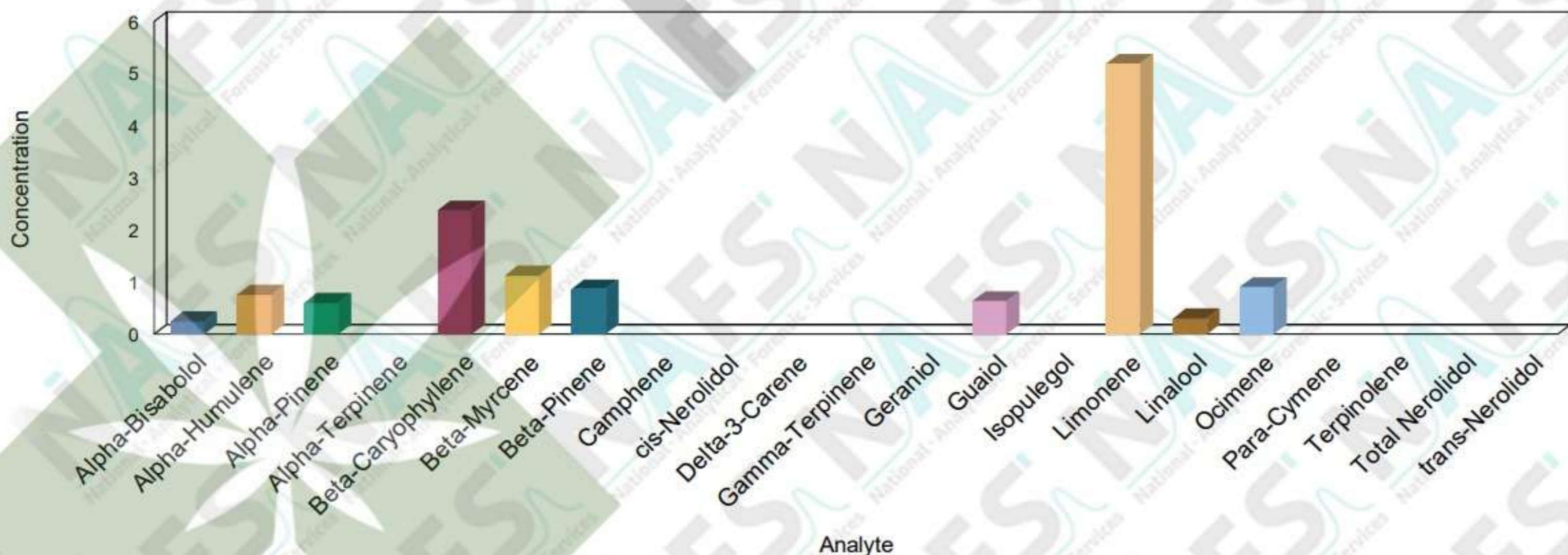
Specification: N/A

Test Method: *TM003.5

Sampling Method: SH007

Analyte	Results	*Std Dev.	*Spec Limits
Alpha-Bisabolol	0.250 mg/g	+/- 0.045 mg/g	N/A
Alpha-Humulene	0.761 mg/g	+/- 0.016 mg/g	N/A
Alpha-Pinene	0.606 mg/g	+/- 0.025 mg/g	N/A
Alpha-Terpinene	ND mg/g	mg/g	N/A
Beta-Caryophyllene	2.383 mg/g	+/- 0.049 mg/g	N/A
Beta-Myrcene	1.131 mg/g	+/- 0.005 mg/g	N/A
Beta-Pinene	0.866 mg/g	+/- 0.040 mg/g	N/A
Camphene	ND mg/g	mg/g	N/A
cis-Nerolidol	ND mg/g	mg/g	N/A
Delta-3-Carene	ND mg/g	mg/g	N/A
Gamma-Terpinene	ND mg/g	mg/g	N/A
Geraniol	ND mg/g	mg/g	N/A
Guaiol	0.638 mg/g	+/- 0.028 mg/g	N/A
Isopulegol	ND mg/g	mg/g	N/A
Limonene	5.198 mg/g	+/- 0.146 mg/g	N/A
Linalool	0.311 mg/g	+/- 0.012 mg/g	N/A
Ocimene	0.905 mg/g	+/- 0.015 mg/g	N/A
Para-Cymene	ND mg/g	mg/g	N/A
Terpinolene	ND mg/g	mg/g	N/A
Total Nerolidol	ND mg/g	mg/g	N/A
trans-Nerolidol	ND mg/g	mg/g	N/A

Analyst: LMARAI 2022/09/12 15:47



*ND (Not Detected)

*The current test method version is Validated

*Standard Deviation (Uncertainty of measurement of applicable duplicate sample)

*Specification Limits (Specified according to pharmacopeial or client specification limits where applicable)

Test Name: TM005.2 (Heavy Metal Class 1,2) (220831-004-001)

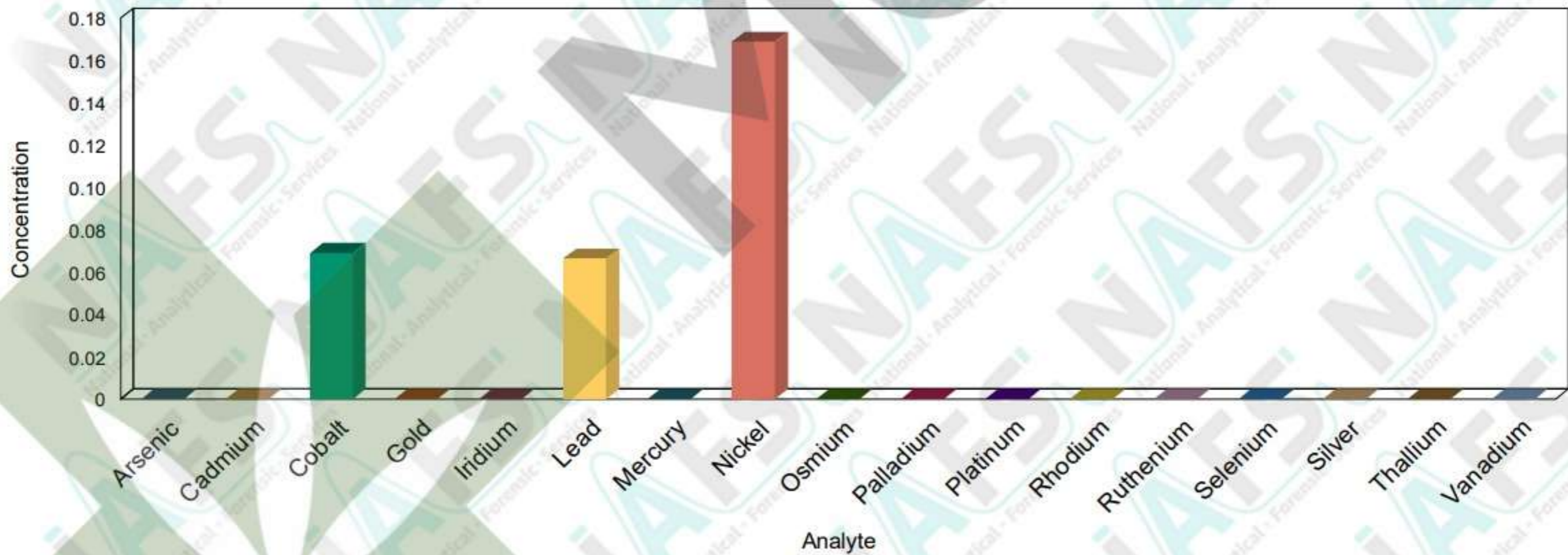
Specification: USP Specification (Inhalation)

Test Method: *TM005.2

Sampling Method: SH007

Analyte	Results	*Std Dev.	*Spec Limits
Silver	ND ppm	ppm	< 0.7 ppm PASS
Arsenic	ND ppm	ppm	< 0.2 ppm PASS
Gold	ND ppm	ppm	< 0.1 ppm PASS
Cadmium	ND ppm	ppm	< 0.2 ppm PASS
Cobalt	0.069 ppm	ppm	< 0.3 ppm PASS
Mercury	ND ppm	ppm	< 0.1 ppm PASS
Iridium	ND ppm	ppm	< 0.1 ppm PASS
Nickel	0.169 ppm OR	+/- 0.001 ppm	< 0.5 ppm PASS
Osmium	ND ppm	ppm	< 0.1 ppm PASS
Lead	0.067 ppm	+/- 0.001 ppm	< 0.5 ppm PASS
Palladium	ND ppm	ppm	< 0.1 ppm PASS
Platinum	ND ppm	ppm	< 0.1 ppm PASS
Rhodium	ND ppm	ppm	< 0.1 ppm PASS
Ruthenium	ND ppm	ppm	< 0.1 ppm PASS
Selenium	ND ppm	ppm	< 13 ppm PASS
Thallium	ND ppm	ppm	< 0.8 ppm PASS
Vanadium	ND ppm	ppm	< 0.1 ppm PASS

Analyst: AWRBKA 2022/09/13 15:08



*ND (Not Detected)

*The current test method version is Validated

*Standard Deviation (Uncertainty of measurement of applicable duplicate sample)

*Specification Limits (Specified according to pharmacopeial or client specification limits where applicable)

*OR (Outside Range) For quantitative analysis, calibration range is the concentration within which the analysis method is accurate. For limit tests the a single point calibration is employed with a range of +/-15% across the specification limit. An OR analyte flag signifies that only semi-quantitative data has been reported.

Interpretations and Opinions:

None

Additions, Deviations & Exclusions:

None

It should be noted that NAFS will only analyze the sample/s received. This sample cannot be regarded as representative of entire batch or crop. It is the responsibility of the client to ensure a representative sample is taken in an appropriate tamper proof sample container. NAFS cannot be held liable for negligent handling, storage and transport of client samples, prior to receipt. Any complaints may be directed toward NAFS using info@nafs.co.za (QA005)

Quality Assurance Manager

Released By: Jeanette Leygonie

Released On: 9/13/2022 3:18:32PM

This certificate of analysis shall not be reproduced except in full, without written approval of NAFS. The information contained in this communication from the sender is confidential. It is intended solely for use by the recipient and others authorized to receive it. If you are not the recipient, you are hereby notified that any disclosure, copying, distribution or taking action in relation to the contents of this information is strictly prohibited and may be unlawful. The results obtained may be for anonymous statistical research purposes. NAFS cannot be held liable for any loss or damage arising from, directly or indirectly related to, sample test results.

Certificate of Analysis

Sample ID: 220831-004-003
Retention Sample ID: 11,978
Client (Code) Name: (CAN0120) MedCan (Pty) Ltd
Client Address: S10 Euro Centre, 363 Rivonia Boulevard, Rivonia
Batch: NA-0005-220822-2
Date Receipt: 2022/09/01 09:45:39
Date Report Released: 2022/09/20 14:11:30
Sample Type: Plant Material
Package Condition: Intact
Received By: JLEYGONIE
Sample Amount: 10 gram
Tamper Proof: No



Test Name: TM009.1 (Total Aerobic Plate Count) (220831-004-003)
Specification: N/A
Test Method: *Outsourced
Sampling Method: SH007

Analyte	Results	*Std Dev.	*Spec Limits
Total Plate Count	0 cfu/g	cfu/g	N/A

Analyst: AWRBKA 2022/09/20 12:21

*ND (Not Detected)
 *The current test method version is Compendial
 *Standard Deviation (Uncertainty of measurement of applicable duplicate sample)
 *Specification Limits (Specified according to pharmacopeial or client specification limits where applicable)

Test Name: TM009.2 (Total Yeast and Mould) (220831-004-003)
Specification: N/A
Test Method: *Outsourced
Sampling Method: SH007

Analyte	Results	*Std Dev.	*Spec Limits
Mould	0 cfu/g	cfu/g	N/A
Total Yeast and Mould	ND cfu/g	cfu/g	N/A
Yeast	0 cfu/g	cfu/g	N/A

Analyst: AWRBKA 2022/09/20 12:21

*ND (Not Detected)
 *The current test method version is Compendial
 *Standard Deviation (Uncertainty of measurement of applicable duplicate sample)
 *Specification Limits (Specified according to pharmacopeial or client specification limits where applicable)

Test Name: TM009.3 (Escherichia Coli) (220831-004-003)

Specification: N/A
Test Method: *Outsourced
Sampling Method: SH007

Analyte	Results	*Std Dev.	*Spec Limits
E. coli	Negative		No Growth PASS

Analyst: AWRBKA 2022/09/20 12:21

*ND (Not Detected)
*The current test method version is Compendial
*Standard Deviation (Uncertainty of measurement of applicable duplicate sample)
*Specification Limits (Specified according to pharmacopeial or client specification limits where applicable)

Test Name: TM009.4 (Salmonella) (220831-004-003)

Specification: N/A
Test Method: *Outsourced
Sampling Method: SH007

Analyte	Results	*Std Dev.	*Spec Limits
Salmonella	Negative		No Growth PASS

Analyst: AWRBKA 2022/09/20 12:21

*ND (Not Detected)
*The current test method version is Compendial
*Standard Deviation (Uncertainty of measurement of applicable duplicate sample)
*Specification Limits (Specified according to pharmacopeial or client specification limits where applicable)

Test Name: TM009.5 (Staphylococcus Aureus) (220831-004-003)

Specification: N/A
Test Method: *Outsourced
Sampling Method: SH007

Analyte	Results	*Std Dev.	*Spec Limits
S. aureus	Negative		No Growth PASS

Analyst: AWRBKA 2022/09/20 12:21

*ND (Not Detected)
*The current test method version is Compendial
*Standard Deviation (Uncertainty of measurement of applicable duplicate sample)
*Specification Limits (Specified according to pharmacopeial or client specification limits where applicable)

Test Name: TM009.6 (Pseudomonas Aeruginosa) (220831-004-003)

Specification: N/A

Test Method: *Outsourced

Sampling Method: SH007

Analyte	Results	*Std Dev.	*Spec Limits
P. aeruginosa	Negative		No Growth PASS

Analyst: AWRBKA 2022/09/20 12:21

*ND (Not Detected)

*The current test method version is Compendial

*Standard Deviation (Uncertainty of measurement of applicable duplicate sample)

*Specification Limits (Specified according to pharmacopeial or client specification limits where applicable)

Interpretations and Opinions:

None

Additions, Deviations & Exclusions:

None

It should be noted that NAFS will only analyze the sample/s received. This sample cannot be regarded as representative of entire batch or crop. It is the responsibility of the client to ensure a representative sample is taken in an appropriate tamper proof sample container. NAFS cannot be held liable for negligent handling, storage and transport of client samples, prior to receipt. Any complaints may be directed toward NAFS using info@nafs.co.za (QA005)

Quality Assurance Manager

Released By: Jeanette Leygonie

Released On: 2022/09/20 14:11:30

This certificate of analysis shall not be reproduced except in full, without written approval of NAFS. The information contained in this communication from the sender is confidential. It is intended solely for use by the recipient and others authorized to receive it. If you are not the recipient, you are hereby notified that any disclosure, copying, distribution or taking action in relation of the contents of this information is strictly prohibited and may be unlawful. The results obtained may be used for anonymous statistical research purposes. NAFS cannot be held liable for any loss or damage arising from, directly or indirectly related to, sample test

**TEST REPORT****MEDCAN (PTY) LTD**
S10 EURO CENTRE, RIVONIA, JOHANNESBURG, S10
EURO CENTRE, RIVONIA, JOHANNESBURG,
JOHANNESBURG, SA**Report No.: 968481****Attn:**

Job/Sample No :	ML-2022-07231-002	Batch/Barcode No :	ML2-0005-210722
Sample Source :	1 (1 BATCH)	Sample Preservation :	Keep at Room Temperature
Container Type :	Original Packaging	Date Received :	25 Jul 2022
Sample Temperature :	25 °C		
Description :	PLANT MATERIAL		
Sample Receipt Temperature :	Acceptable Condition		

Pharmaceuticals Date Reported:02/08/2022

Test	Method	Units	LOQ	Results	Guideline / Specification limit
					Aflatoxin - Peanuts
Aflatoxin B1	HP040-56-W (Based on the Vicam Aflatest WP Method)	µg/kg	0.26	Not Detected	NMT 5µg/kg
Total Aflatoxins	HP040-56-W (Based on the Vicam Aflatest WP Method)	µg/kg	0.86	Not Detected	NMT 10µg/kg

Tests marked with an () indicate that those tests are not SANAS Accredited and are not included in the SANAS Schedule of Accreditation*

<10 / <1= Not detected / less than the lower detection limit of the test method, for the specified sample type / volume of sample tested

Remarks:This is to certify that the above mentioned sample is conforming to Aflatoxin - Peanuts in terms of the above tested Pharmaceuticals

Reported by**Nazma Ismail**



M&L LABORATORY SERVICES (PTY) LTD.

Pesticide Residue Screen on one Cannabis Sample

Client Details:

MedCan (PTY) LTD
S10 Euro Centre, Rivonia
Johannesburg
Attention: Dumani Dlamini

Testing Facility:

M and L Laboratory Services (Pty) Ltd
40 Modulus Road, Ormonde,
Johannesburg, 2091



Certificate



All services are rendered in accordance with Bureau Veritas M&L Laboratory Services (Pty) Ltd General Terms and conditions of Business, which has been supplied to you, this certificate cannot be reproduced except in full without the written consent of M and L Laboratory Services.



M&L LABORATORY SERVICES (PTY) LTD.

1. Method :

Pesticides Screening by UPLC-MS-MS and GC-MS-MS based on **PH Eur 2.8.13** method

2. Sample Description and Details:

2.1. Sample Description

Sample Name.	Plant Material: Cannabis Flower
Supplier	MedCan
Batch:	ML1-0005-210722
Lab No.	968480

Pesticide Detected	Residue mg/kg
--------------------	---------------

None Detected	
---------------	--

Authorised Signature

Certificate



MedCan

All services are rendered in accordance with Bureau Veritas M&L Laboratory Services (Pty) Ltd General Terms and conditions of Business, which has been supplied to you, this certificate cannot be reproduced except in full without the written consent of M and L Laboratory Services.

M&L LABORATORY SERVICES (PTY) LTD.

The samples were screened for the following pesticides below and above were detected above their Reporting limit mg/kg shown below, unless specified in the body of the report.

Pesticides Screened below by GC-MS and LC-MS:

LIST 1

Pesticides Compounds	Reporting Limits (mg/kg)	Pesticides Compounds	Reporting Limits (mg/kg)	Pesticides Compounds	Reporting Limits (mg/kg)
2,4-D	0.010	Demeton-S-methyl-sulfone	0.010	Guazatine	0.010
Acetamiprid	0.010	Desmethyl-pirimicarb	0.010	Haloxypop R methyl	0.010
Aclonifen	0.010	Diafenthuron	0.010	Heptachlor	0.010
Acrinathrin	0.010	Diazinon	0.010	Hexachlorbenzene	0.010
Acephate	0.010	Dichlofluanid	0.010	Hexachlorocyclohexane	0.010
Alachlor	0.010	Dichlorvos	0.010	Imazalil	0.010
Aldrin and dieldrin	0.010	Diclofop methyl	0.010	Imidacloprid	0.010
Ametryn	0.010	Dicofol	0.010	Indoxacarb	0.010
Amitraz	0.010	Dieldrin	0.010	Iprodione	0.010
Atrazine	0.010	Diethofencarb	0.010	Iprovalicarb	0.010
AvermectinB1a	0.010	Difenoconazole	0.010	Kresoxim methyl	0.010
AvermectinB1b	0.010	Diflubenazuron	0.010	Lambda cyhalothrin	0.010
Azinphos ethyl	0.010	Diflubenazuron	0.010	Lindan	0.010
Azinphos methyl	0.010	Diflufenican	0.010	Linuron	0.010
Azoxystrobin	0.010	Dimethenamid	0.010	Lufenuron	0.010
Benalaxyl	0.010	Dimethoate	0.010	Malaoxon	0.010
Benfluralin	0.010	Dimethomorph	0.010	Malathion	0.010
Benfuracarb	0.010	Diniconazole	0.010	Mandipropamide	0.010
Bentazone	0.010	Diphenylamine	0.010	Maneb	0.010
Bifenazate	0.010	Diuron	0.010	Mepanpyrim	0.010
Bifenthrin	0.010	Dithiocarbamates	0.010	Meptyldinocap	0.010
Bitertanol	0.010	Emamectin benzoate B1a	0.010	Metalaxyl-M	0.010
Boscalid	0.010	Emamectin B1b	0.010	Metamitron	0.010
Bromacil	0.010	Endosulfan	0.010	Mecarbam	0.010
Bromide, inorganic	0.010	Endrin	0.010	Methacriphos	0.010
Bromophos ethyl	0.010	Endrin aldehyde	0.010	Methamidophos	0.010
Bromophos methyl	0.010	Esfenvalerate	0.010	Methidathion	0.010
Bromuconazole	0.010	Ethalfuralin	0.010	Methiocarb	0.010
Bromopropylate	0.010	Ethoxyquin	0.010	Methomyl	0.010
Bupimate	0.010	Ethion	0.010	Methoxyfenozide	0.010
Buprofezin	0.010	Etofenprox	0.010	Metolachlor-S	0.010
Cadusafos	0.010	Etoazole	0.010	Metominostrobin	0.010
Captan	0.010	Etrimphos	0.010	Methoxychlor	0.010
Carbaryl	0.010	Fenamidone	0.010	Metoconazole - cis	0.010
Carbendazim	0.010	Fenamiphos	0.010	Metoconazole - trans	0.010
Carbosulfan	0.010	Fenarimol	0.010	Metribuzin	0.010
Carbofuran	0.010	Fenzaquin	0.010	Myclobutanil	0.010
Carbofuran, 3-hydroxy	0.010	Fenbuconazole	0.010	Mirex	0.010
Carfentrazone ethyl	0.010	Fenhexamide	0.010	Monocrotophos	0.010
Chlorantraniliprole	0.010	Fenoxycarb	0.010	Noovaluron	0.010
Chlorfenapyr	0.010	Fenchlorophos	0.010	Omethoate	0.010
Chlorfluazuron	0.010	Fenitrothion	0.010	Ortho phenylphenol	0.010
Chlorothalonil	0.010	Fenoxaprop-p-ethyl	0.010	Oxadiargyl	0.010
Chlorpropham	0.010	Fenpropathrin	0.010	Oxadiazon	0.010
Chlordane	0.010	Fenpyroximate	0.010	Oxamyl	0.010
Chlorfenvinphos	0.010	Fensulfothion	0.010	Oxydemeton methyl	0.010



M&L LABORATORY SERVICES (PTY) LTD.

Chlorpyrifos - ethyl	0.010	Fenthion	0.010	Oxyfluorfen	0.010
Chlorpyrifos - methyl	0.010	Fenvalerate	0.010	Paclobutrazol	0.010
Chlorthal - dimethyl	0.010	Fluazifop-p-butyl	0.010	Parathion ethyl	0.010
Clofentezine	0.010	Fluazinam	0.010	Parathion methyl	0.010
Clomazone	0.010	Fludioxinil	0.010	Penconazole	0.010
Clothiandin	0.010	Flucytrinate	0.010	Pentachloranisole	0.010
Cyantraniliprole	0.010	Flufenoxuron	0.010	Pendimethalin	0.010
Cycloxydim	0.010	Fluopicolide	0.010	Phosalone	0.010
Cyflufenamid	0.010	Flurochloridone	0.010	Phosmet	0.010
Cyfluthrin (1, 2, 3, & 4)	0.010	Fluroxypyr meptyl	0.010	Phosmet-oxon	0.010
Cymoxanil	0.010	Fluvalinate & tau	0.010	Piperonyl butoxide	0.010
Cypermethrin	0.010	Fonophos	0.010	Pirimicarb	0.010
Cyproconazole	0.010	Fusilazole	0.010	Pirimiphos-methyl	0.010
Cyprodinil	0.010	Flutolanil	0.010	Prochloraz	0.010
Cyhalothrin	0.010	Flutriafol	0.010	Phosmet-oxon	0.010
DDT, DDE & TDE	0.010	Folpet	0.010	Piperonyl butoxide	0.010
Deltamethrin	0.010	Formetanate HCl	0.010	Pirimicarb	0.010

Pesticides Compounds	Reporting Limits (mg/kg)	Pesticides Compounds	Reporting Limits (mg/kg)	Pesticides Compounds	Reporting Limits (mg/kg)
Pirimiphos-methyl	0.010	Quinalphos	0.010	Tetradifon	0.010
Prochloraz	0.010	Quintozene	0.010	Thiacloprid	0.010
Procymidone	0.010	Rimsulfuron	0.010	Thiamethoxam	0.010
Profenophos	0.010	S-421	0.010	Thiodicarb	0.010
Prometryne	0.010	Simazine	0.010	Thiophanate methyl	0.010
Propamocarb HCl	0.010	Spinetoram J	0.010	Tolclophos methyl	0.010
Propargate	0.010	Spineteram L	0.010	Triadimefon	0.010
Propyzamide	0.010	Sulfoxaflor	0.010	Triadimenol	0.010
Proquinazid	0.010	Spinosyn A	0.010	Triasulfuron	0.010
Prothiosphos	0.010	Spinosyn B	0.010	Trifloxystrobin	0.010
Pymetrozine	0.010	Spirodiclofen	0.010	Triflumuron	0.010
Pyraclostrobin	0.010	Spiromesifen	0.010	Trifluraline	0.010
Pyraflufen ethyl	0.010	Spiroxamine	0.010	Uniconazole	0.010
Pyridaben	0.010	Sulfoxaflor	0.010	Vinclozolin	0.010
Pyridalyl	0.010	Tebuconazole	0.010	Zoxamide	0.010
Pyrimethanil	0.010	Tebufenpyrad	0.010	Permethrin isomers	0.010
Pyrimidifen	0.010	Tecnazene	0.010	Acetochlor	0.010
Pyriproxyfen	0.010	Teflubenzuron	0.010	Terbuthylazine	0.010
Pyrethrum	0.010	Terbacil	0.010		
Quinoxifen	0.010	Terbutryne	0.010		



M&L LABORATORY SERVICES (PTY) LTD.
EXTRACT OF THE BUREAU VERITAS GENERAL TERMS AND CONDITIONS OF BUSINESS

This extract of the Bureau Veritas general terms and conditions of business ("**General Conditions**") shall govern all services, including (but not limited to) laboratory test work, surveys, sampling, site investigations, consultations and opinions, performed for any individual or juristic person (the "**Client**") by M and L Laboratory Services Proprietary Limited, its subsidiary companies and their employees, agents, consultants and subcontractors (collectively referred to as the "**Company**"), whether in terms of a specified contract or not. For the purpose of these General Conditions, the Company and the Client shall collectively be referred to as the "**Parties**" and individually as a "**Party**".

1. QUOTATIONS

Any quotations for Services submitted by the Company to the Client shall be based on information supplied to the Company by the Client and will not under any circumstances be binding on the Company if such information is incorrect or incomplete in any manner.

2. INSTRUCTIONS

The Client will provide the Company with clear and precise written instructions, documents, information and samples prior to the performance of the Services. The Company will not be liable for any error, omission or inaccuracy in the reports or certificates produced by it to the extent that the Company has been given erroneous or incomplete information by the Client. The reports and certificates produced by the Company reflect the findings of the Company at the time of performance of the Services only.

3. SAMPLE MATERIAL

The Client will ensure that all samples/materials submitted by it for test work of any nature are clearly marked and identifiable. Should it be necessary for the Company to carry out any sample preparation, preliminary experimental work, or research prior to carrying out the Services, the Client will be liable for any charges in respect thereof. Unless the Client otherwise instructs in writing, the Company may retain, return to the Client, destroy or dispose of all excess samples, material, specimens or exhibits provided by the Client to the Company as soon as the Services are completed and the results have been reported to the Client. Any destruction or disposal shall exclude normal amounts of reserve sample material which the Company shall retain for a period of three months from date of completion of the Services. If the Client requires the Company to return any samples or materials to it or a third party, all costs associated therewith, including associated telecommunication costs, will be borne and paid for the Client.

4. FEES AND TERMS OF PAYMENT

In consideration for the provision of the Services by the Company, the Client shall pay the fees calculated in accordance with the Company's tariff of fees at the time, copies of which may be requested by the Client at any time. In the event of any changes in the Company's fees, the Company shall provide written notification thereof to the Client within a reasonable time prior to such new fees becoming effective. If the Client does not have an account with the Company, the Client shall be required to pay the whole or part of the fees before the Company will commence the Services or release the results, as the case may be. The Client will pay each valid invoice submitted to it by the Company in full and in cleared funds within 30 days of the date of the invoice. The Company shall be entitled to charge interest at 2% per month on any amounts not paid on the due date.

5. LIABILITY AND INDEMNITY

Neither Party shall be liable to the other Party for any consequential, indirect, incidental or special losses or damages of any nature whatsoever and howsoever arising. Without prejudice to the foregoing, the total liability of the Company arising out of or in connection with this Agreement or in relation to the Services shall be limited to the fee paid or payable by the Client to the Company for the Services that gave rise to the Company's liability to the Client, if any. The Client indemnifies the Company and holds it harmless against all claims made by third parties for losses, damages or expenses of whatsoever nature and howsoever arising relating to the performance, purported performance or non-performance of any Services to the extent that the aggregate of such claims for any one Service exceeds the limitation of liability set out in this clause 5.

6. PROVISION OF THE SERVICES

The Company shall provide the Services with reasonable care, skill and diligence as expected of a competent body experienced in performing services of a similar nature and under similar circumstances. If the Client is aware of any apparent inaccuracy in any results reported by the Company in respect of the Services, the Client shall immediately advise the Company accordingly, and allow the Company a reasonable opportunity to check such results and amend them if necessary.

7. PUBLICATION OF RESULTS

M&L LABORATORY SERVICES (PTY) LTD.

Any reports or certificates issued by the Company are intended for the exclusive use of the Client and shall not be published, used for advertising purposes, copied or replicated for distribution to any person or entity or otherwise publicly disclosed without the prior written consent of the Company.

8. ALTERATIONS OF TERMS

No employee, agent or representative of the Company is authorised to alter or waive any of the terms contained in these General Conditions unless in writing and signed by or on behalf of the Parties. The performance of any test shall further be subject to any additional special conditions as the Company may impose from time to time. If such special conditions differ from any provisions set out herein, such special conditions shall, to the extent of such difference, take precedence.

9. LAW OF SOUTH AFRICA

These General Conditions shall be governed by and construed in accordance with the laws of the Republic of South Africa. The Parties irrevocably consent to the jurisdiction of the South Gauteng High Court, Johannesburg, if any dispute or claim arises out of or in connection with this Agreement.

For full business terms and conditions please click or visit http://portal.bureauveritas.co.za/downloads/conditions_ml.pdf

MediScan





For more information, please contact
info@medcan.co.za