



AltMed Arizona - Verano AZC  
 1341 W. Industrial Dr.  
 Coolidge, AZ 85128  
 License #: 00000105DCOU00194638  
 Sample ID: 2402SMAZ0282.0885  
 Batch #: 240221BGGG



**CERTIFICATE OF ANALYSIS**  
 License #: 00000020LCVT89602592

Certificate: 4477

## BITS Guava GO

Batch #: 240221BGGG  
 Strain: Sativa Blend  
 Parent Batch #: 231121MDIS  
 Production Method: Alcohol  
 Harvest Date: 11/21/2023  
 Received: 02/26/2024

Sample ID: 2402SMAZ0282.0885  
 Amount Received: 56.5 g  
 Sample Type: Soft Chew  
 Sample Collected: 02/26/2024 11:06:00  
 Manufacture Date: 02/21/2024  
 Published: 02/28/2024



## COMPLIANCE FOR RETAIL

### Regulated Analytes

Cannabinoid Profile (Q3) <b>Tested</b>	Microbial Contaminants <b>Pass</b>	Residual Solvents <b>Not Tested</b>
Pesticides, Fungicides, and Growth Regulators <b>Not Tested</b>	Mycotoxins <b>Not Tested</b>	Heavy Metals <b>Not Tested</b>

### Additional Analytes (Not Regulated)

Terpenes Total (Q3) <b>Not Tested</b>	Moisture Analysis (Q3) <b>Not Tested</b>	Water Activity (Q3) <b>Not Tested</b>
Filth & Foreign (Q3) <b>Not Tested</b>	Homogeneity (Q3) <b>Not Tested</b>	Additional Microbial Contaminants (Q3) <b>Not Tested</b>

<b>0.208%</b> Total THC
<b>&lt;LOQ</b> Total CBD
<b>&lt;LOQ</b> CBN
<b>0.007%</b> CBG
<b>0.218%</b> Total Cannabinoids (Q3)

Ahmed Munshi  
 Technical Laboratory Director

Smithers CTS Arizona LLC  
 734 W Highland Avenue, 2nd Floor  
 Phoenix, AZ 85013  
 (602) 806-6930



The product associated with this COA has been tested by Smithers CTS Arizona LLC, using validated state certified testing methodologies as required by Arizona state law. Testing results were obtained according to Smithers' quality assurance plan and requirements found in R9-17-404.03 and R9-17-404.04. This COA is governed by the terms and conditions listed on: <https://www.smithers.com/arizona-terms-conditions>



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**Cannabinoid Profile**

HPLC

Tested

**Sample Prep**

Batch Date: 02/27/2024  
 SOP: 418.AZ  
 Batch Number: 968

**Sample Analysis**

Date: 02/28/2024  
 SOP: 417.AZ - HPLC  
 Sample Weight: 1.002 g  
 Volume: 10 mL

Analyte	LOD (mg/g)	LOQ (mg/g)	Dil.	Actual % (w/w)	mg/g	mg/serving	mg/package	Qualifier
CBC	0.003	0.010	1	0.002	0.020	0.057	1.132	M3
CBD	0.003	0.010	1	<LOQ	<LOQ	<LOQ	<LOQ	M3
CBDA	0.003	0.010	1	ND	ND	ND	ND	M3
CBDV	0.003	0.010	1	ND	ND	ND	ND	M3
CBG	0.003	0.010	1	0.007	0.069	0.195	3.905	M3
CBGA	0.003	0.010	1	ND	ND	ND	ND	M3
CBN	0.003	0.010	1	<LOQ	<LOQ	<LOQ	<LOQ	M3
d8-THC	0.003	0.010	1	ND	ND	ND	ND	M3
d9-THC	0.003	0.010	1	0.208	2.083	5.895	117.898	M3
THCA	0.003	0.010	1	ND	ND	ND	ND	M3
THCV	0.003	0.010	1	0.001	0.014	0.040	0.792	M3

Cannabinoid Totals	Actual % (w/w)	mg/g	mg/serving	mg/package	Qualifier
Total THC	0.208	2.083	5.895	117.898	
Total CBD	<LOQ	<LOQ	<LOQ	<LOQ	
Total Cannabinoids	0.218	2.185	6.184	123.671	Q3

Total THC = THC + (0.877 x THCA) and Total CBD = CBD + (0.877 x CBDA)  
 ND = Not Detected, NT = Not Tested, <LOQ = Below Limit of Quantitation  
 Serving Weight: 2.83 None; Servings/Package: 20

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Certificate: 4477

**Microbial Analysis**  
 Pass

**Sample Prep**

Batch Date: 02/27/2024  
 SOP: 431.AZ  
 Batch Number: 967

**Sample Analysis**

Date: 02/28/2024  
 SOP: 431.AZ - TEMPO (MPN)  
 Sample Weight: 1.095 g

Analyte	Allowable Criteria	Actual Result	Pass/Fail	Qualifier
E. coli	< 10 CFU/g	< 10 CFU/g	Pass	

**Sample Prep**

Batch Date: 02/27/2024  
 SOP: 406.AZ  
 Batch Number: 972

**Sample Analysis**

Date: 02/28/2024  
 SOP: 406.AZ - qPCR (MG)  
 Sample Weight: 1.007 g

Analyte	Allowable Criteria	Actual Result	Pass/Fail	Qualifier
Salmonella	Not Detected in One Gram	Not Detected in One Gram	Pass	

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 Batch #: 240221BGGG



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## Qualifier Legend

- B1** The target analyte detected in the calibration is at or above the limit of quantitation, but the sample result for potency testing, is below the limit of quantitation.
- B2** The target analyte detected in the calibration blank, or the method blank is at or above the limit of quantitation, but the sample result when testing for pesticides, fungicides, herbicides, growth regulators, heavy metals, or residual solvents, is below the maximum allowable concentration for the analyte.
- D1** The limit of quantitation and the sample results were adjusted to reflect sample dilution.
- I1** The relative intensity of a characteristic ion in a sample analyte exceeded the acceptance with respect to the reference spectra, indicating interference.
- L1** When testing for pesticides, fungicides, herbicides, growth regulators, heavy metals, or residual solvents, the percent recovery of a laboratory control sample is greater than the acceptance limits, but the sample's target analytes were not detected above the maximum allowable concentrations for the analytes in the sample.
- M1** The recovery from the matrix spike was high, but the recovery from the laboratory control sample was within acceptance criteria.
- M2** The recovery from the matrix spike was low, but the recovery from the laboratory control sample was within acceptance criteria.
- M3** The recovery from the matrix spike was unusable because the analyte concentration was disproportionate to the spike level, but the recovery from the laboratory control sample was within acceptance criteria.
- M4** The analysis of a spiked sample required a dilution such that the spike recovery calculation does not provide useful information, but the recovery from the associated laboratory control sample was within acceptance criteria.
- M5** The analyte concentration was determined by the method of standard addition, in which the standard is added directly to the aliquots of the analyzed sample.
- M6** A description of the variance is described in the final report of testing according to R9-17- 404.06(B)(3)(d)(ii).
- Q1** Sample integrity was not maintained.
- Q2** The sample is heterogeneous, and sample homogeneity could not be readily achieved using routine laboratory practices.
- Q3** Testing result is for informational purposes only and cannot be used to satisfy dispensary testing requirements in R9-17-317.01(A) or labeling requirements in R9-17-317.
- R1** The relative percent difference for the laboratory control sample and duplicate exceeded the limit, but the recovery was within acceptance criteria.
- R2** The relative percent difference for a sample and duplicate exceeded the limit.
- V1** The recovery from continuing calibration verification standards exceeded the acceptance limits, but the sample's target analytes were not detected above the maximum allowable for the analytes in the sample.

**Cultivated By:** Fort Consulting, Llc 00000105DCOU00194638/00000064ESAK09838873

**Manufactured By:** Fort Consulting, Llc 00000105DCOU00194638/00000064ESAK09838873

**Disclaimer:** Using marijuana during pregnancy could cause birth defects or other health issues to your unborn child.

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**Notes:** Expiration Date: 02/21/2025  
 Date Sample Prepared: 02/21/2024



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Accreditation #: 103104

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 License #: 00000105DCOU00194638  
 Sample ID: 2311SMAZ1800.7047  
 Batch #: 231121MDIS



**CERTIFICATE OF ANALYSIS**  
 License #: 00000020LCVT89602592

Certificate: 2308

## Distillate

Batch #: 231121MDIS      Sample ID: 2311SMAZ1800.7047  
 Strain: Hybrid Blend      Amount Received: 7.4 g  
 Parent Batch #:      Sample Type: Distillate  
 Sample Collected: 11/29/2023 12:00:00      Received: 11/29/2023  
 Published: 12/06/2023



## COMPLIANCE FOR RETAIL

### Regulated Analytes

Cannabinoid Profile (Q3) <b>Tested</b>	Microbial Contaminants <b>Pass</b>	Residual Solvents <b>Pass</b>
Pesticides, Fungicides, and Growth Regulators <b>Pass</b>	Mycotoxins <b>Pass</b>	Heavy Metals <b>Pass</b>

### Additional Analytes (Not Regulated)

Terpenes Total (Q3) <b>Not Tested</b>	Moisture Analysis (Q3) <b>Not Tested</b>	Water Activity (Q3) <b>Not Tested</b>
Filth & Foreign (Q3) <b>Not Tested</b>	Homogeneity (Q3) <b>Not Tested</b>	Additional Microbial Contaminants (Q3) <b>Not Tested</b>

<b>92.356%</b> Total THC
<b>0.274%</b> Total CBD
<b>0.279%</b> CBN
<b>3.106%</b> CBG
<b>97.569%</b> Total Cannabinoids (Q3)

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Certificate: 2308

**Cannabinoid Profile**

HPLC                      **Tested**

**Sample Prep**  
 Batch Date: 11/29/2023  
 SOP: 418.AZ  
 Batch Number: 444

**Sample Analysis**  
 Date: 12/06/2023  
 SOP: 417.AZ - HPLC  
 Sample Weight: 0.040 g  
 Volume: 40 mL

Analyte	LOD (mg/g)	LOQ (mg/g)	Dil.	Actual % (w/w)	mg/g	Qualifier
CBC	0.644	1.954	2	0.899	8.992	
CBD	0.644	1.954	2	0.274	2.744	
CBDA	0.644	1.954	2	ND	ND	
CBDV	0.644	1.954	2	ND	ND	
CBG	0.644	1.954	2	3.106	31.062	
CBGA	0.644	1.954	2	ND	ND	
CBN	0.644	1.954	2	0.279	2.788	
d8-THC	0.644	1.954	2	ND	ND	
d9-THC	0.644	1.954	2	92.356	923.556	
THCA	0.644	1.954	2	ND	ND	
THCV	0.644	1.954	2	0.655	6.546	

Cannabinoid Totals	Actual % (w/w)	mg/g	Qualifier
Total THC	92.356	923.556	
Total CBD	0.274	2.744	
Total Cannabinoids	97.569	975.688	Q3

Total THC = THC + (0.877 x THCA) and Total CBD = CBD + (0.877 x CBDA)  
 ND = Not Detected, NT = Not Tested, <LOQ = Below Limit of Quantitation

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Certificate: 2308

**Microbial Analysis**  
**Pass**

**Sample Prep**

Batch Date: 11/30/2023  
 SOP: 431.AZ  
 Batch Number: 449

**Sample Analysis**

Date: 12/06/2023  
 SOP: 431.AZ - TEMPO (MPN)  
 Sample Weight: 1.006 g

Analyte	Allowable Criteria	Actual Result	Pass/Fail	Qualifier
E. coli	< 100 CFU/g	< 10 CFU/g	Pass	

**Sample Prep**

Batch Date: 11/30/2023  
 SOP: 406.AZ  
 Batch Number: 448

**Sample Analysis**

Date: 12/06/2023  
 SOP: 406.AZ - qPCR (MG)  
 Sample Weight: 1.009 g

Analyte	Allowable Criteria	Actual Result	Pass/Fail	Qualifier
Salmonella	Not Detected in One Gram	Not Detected in One Gram	Pass	

**Sample Prep**

Batch Date: 11/30/2023  
 SOP: 406.AZ  
 Batch Number: 448

**Sample Analysis**

Date: 12/06/2023  
 SOP: 406.AZ - qPCR (MG)  
 Sample Weight: 1.009 g

Analyte	Allowable Criteria	Actual Result	Pass/Fail	Qualifier
Aspergillus flavus	Not Detected in One Gram	Not Detected in One Gram	Pass	
Aspergillus fumigatus	Not Detected in One Gram	Not Detected in One Gram	Pass	
Aspergillus niger	Not Detected in One Gram	Not Detected in One Gram	Pass	
Aspergillus terreus	Not Detected in One Gram	Not Detected in One Gram	Pass	

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**Residual Solvents**

HS-GC-MS                      **Pass**

**Sample Prep**

Batch Date: 12/01/2023  
 SOP: 405.AZ  
 Batch Number: 452

**Sample Analysis**

Date: 12/06/2023  
 SOP: 405.AZ - HS-GC-MS  
 Sample Weight: 0.052 g

Analyte	LOD / LOQ (ppm)	Dil.	Action Limit (ppm)	Results (ppm)	Qualifier	Analyte	LOD / LOQ (ppm)	Dil.	Action Limit (ppm)	Results (ppm)	Qualifier
Acetone	63 / 192	1	1000	ND		Heptane	321 / 962	1	5000	ND	
Acetonitrile	27 / 79	1	410	ND		Hexanes	46 / 139	1	290	ND	
Benzene	0.13 / 0.38	1	2	ND		Isopropyl acetate	321 / 962	1	5000	ND	
Butanes	160 / 481	1	5000	ND		Methanol	192 / 577	1	3000	ND	
Chloroform	4 / 12	1	60	ND		Pentanes	321 / 962	1	5000	ND	
Dichloromethane	38 / 115	1	600	ND		2-Propanol (IPA)	321 / 962	1	5000	ND	
Ethanol	321 / 962	1	5000	ND		Toluene	58 / 171	1	890	ND	
Ethyl acetate	321 / 962	1	5000	ND		Xylenes	279 / 835	1	2170	ND	
Ethyl ether	321 / 962	1	5000	ND							

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**Heavy Metals**

ICP-MS **Pass**

**Sample Prep**  
 Batch Date: 12/01/2023  
 SOP: 428.AZ  
 Batch Number: 454

**Sample Analysis**  
 Date: 12/06/2023  
 SOP: 428.AZ - ICP-MS  
 Sample Weight: 0.219 g  
 Volume: 6 mL

Analyte	LOD (ppm)	LOQ (ppm)	Dil.	Action Limit (ppm)	Results (ppm)	Qualifier
Arsenic	0.018	0.183	10	0.4	ND	
Cadmium	0.018	0.183	10	0.4	ND	
Lead	0.018	0.457	10	1	ND	
Mercury	0.018	0.091	10	0.2	ND	

**Mycotoxin Analysis**

LC-MS/MS **Pass**

**Sample Prep**  
 Batch Date: 11/29/2023  
 SOP: 432.AZ  
 Batch Number: 446

**Sample Analysis**  
 Date: 12/06/2023  
 SOP: 424.AZ - LC-MS/MS  
 Sample Weight: 0.556 g  
 Volume: 12.5 mL

Analyte	LOD (ppb)	LOQ (ppb)	Dil.	Action Limit (ppb)	Results (ppb)	Qualifier
Total Aflatoxins	3.60	8.99	1	20	ND	I1, M2V1
Aflatoxin B1	3.60	8.99	1	0	ND	I1, M2V1
Aflatoxin B2	3.60	8.99	1	0	ND	V1
Aflatoxin G1	3.60	8.99	1	0	ND	I1, V1
Aflatoxin G2	3.60	4.50	1	0	ND	V1
Ochratoxin A	8.99	8.99	1	20	ND	I1

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**Pesticides, Fungicides, and  
 Growth Regulators**  
 LC-MS/MS **Pass**

**Sample Prep**

Batch Date: 11/29/2023  
 SOP: 432.AZ  
 Batch Number: 446

**Sample Analysis**

Date: 12/06/2023  
 SOP: 424.AZ - LC-MS/MS  
 Sample Weight: 0.556 g  
 Volume: 12.5 mL

Analyte	LOD / LOQ (ppm)	Dil.	Action Limit (ppm)	Results (ppm)	Qualifier	Analyte	LOD / LOQ (ppm)	Dil.	Action Limit (ppm)	Results (ppm)	Qualifier
Abamectin B1a	0.075 / 0.225	1	0.5	ND	M2 V1	Hexythiazox	0.150 / 0.450	1	1	ND	M2
Acephate	0.060 / 0.180	1	0.4	ND	V1	Imazalil	0.030 / 0.090	1	0.2	ND	M2 V1
Acetamiprid	0.030 / 0.090	1	0.2	ND	V1	Imidacloprid	0.060 / 0.180	1	0.4	ND	V1
Aldicarb	0.060 / 0.180	1	0.4	ND		Kresoxim-methyl	0.060 / 0.180	1	0.4	ND	M2 V1
Azoxystrobin	0.030 / 0.090	1	0.2	ND	V1	Malathion	0.030 / 0.090	1	0.2	ND	V1
Bifenazate	0.030 / 0.090	1	0.2	ND		Metaxalyl	0.030 / 0.090	1	0.2	ND	V1
Bifenthrin	0.030 / 0.090	1	0.2	<LOQ	V1	Methiocarb	0.030 / 0.090	1	0.2	ND	V1
Boscalid	0.060 / 0.180	1	0.4	ND	M2 V1	Methomyl	0.060 / 0.180	1	0.4	ND	V1
Carbaryl	0.030 / 0.090	1	0.2	ND	V1	Myclobutanil	0.030 / 0.090	1	0.2	ND	V1
Carbofuran	0.030 / 0.090	1	0.2	ND	V1	Naled	0.075 / 0.225	1	0.5	ND	M2 V1
Chlorantraniliprole	0.030 / 0.090	1	0.2	ND	V1	Oxamyl	0.150 / 0.450	1	1	ND	V1
Chlorfenapyr	0.150 / 0.450	1	1	ND	I1, M2 R1	Paclobutrazol	0.060 / 0.180	1	0.4	ND	M2 V1
Chlorpyrifos	0.030 / 0.090	1	0.2	ND	M2	Permethrins	0.030 / 0.090	1	0.2	ND	V1
Clofentezine	0.030 / 0.090	1	0.2	ND	M2 V1	Phosmet	0.030 / 0.090	1	0.2	ND	M2 V1
Cyfluthrin	0.150 / 0.450	1	1	ND	M2 V1	Piperonyl Butoxide	0.299 / 0.899	1	2	ND	V1
Cypermethrin	0.150 / 0.450	1	1	ND	M2 V1	Prallethrin	0.030 / 0.090	1	0.2	ND	V1
Daminozide	0.150 / 0.450	1	1	ND		Propiconazole	0.060 / 0.180	1	0.4	ND	M2 V1
Diazinon	0.030 / 0.090	1	0.2	ND	V1	Propoxur	0.030 / 0.090	1	0.2	ND	V1
Dichlorvos	0.015 / 0.045	1	0.1	ND	M2 V1	Pyrethrins	0.126 / 0.377	1	1	ND	V1
Dimethoate	0.030 / 0.090	1	0.2	ND	V1	Pyridaben	0.030 / 0.090	1	0.2	ND	V1
Ethoprophos	0.030 / 0.090	1	0.2	ND	V1	Spinosad	0.030 / 0.090	1	0.2	ND	M2 V1
Etofenprox	0.060 / 0.180	1	0.4	ND	V1	Spiromesifen	0.030 / 0.090	1	0.2	ND	V1
Etoxazole	0.030 / 0.090	1	0.2	ND	V1	Spirotetramat	0.030 / 0.090	1	0.2	ND	V1
Fenoxycarb	0.030 / 0.090	1	0.2	ND	V1	Spiroxamine	0.060 / 0.180	1	0.4	ND	V1
Fenpyroximate	0.060 / 0.180	1	0.4	ND	V1	Tebuconazole	0.060 / 0.180	1	0.4	ND	V1
Fipronil	0.060 / 0.180	1	0.4	ND	V1	Thiacloprid	0.030 / 0.090	1	0.2	ND	V1
Fonicamid	0.150 / 0.450	1	1	ND	V1	Thiamethoxam	0.030 / 0.090	1	0.2	ND	V1
Fludioxonil	0.060 / 0.180	1	0.4	ND	M2 V1	Trifloxystrobin	0.030 / 0.090	1	0.2	ND	M2 V1

**Ahmed Munshi**  
 Technical Laboratory Director

*Ahmed Munshi*

**Smithers CTS Arizona LLC**  
 734 W Highland Avenue, 2nd Floor  
 Phoenix, AZ 85013  
 (602) 806-6930



The product associated with this COA has been tested by Smithers CTS Arizona LLC, using validated state certified testing methodologies as required by Arizona state law. This COA is governed by the terms and conditions listed on: <https://www.smithers.com/arizona-terms-conditions>



AltMed Arizona - Verano AZC  
 1341 W. Industrial Dr.  
 Coolidge, AZ 85128  
 License #: 00000105DCOU00194638  
 Sample ID: 2311SMAZ1800.7047  
 Batch #: 231121MDIS



**CERTIFICATE OF ANALYSIS**  
 License #: 00000020LCVT89602592

Certificate: 2308

**Qualifier Legend**

- B1** The target analyte detected in the calibration is at or above the limit of quantitation, but the sample result for potency testing, is below the limit of quantitation.
- B2** The target analyte detected in the calibration blank, or the method blank is at or above the limit of quantitation, but the sample result when testing for pesticides, fungicides, herbicides, growth regulators, heavy metals, or residual solvents, is below the maximum allowable concentration for the analyte.
- D1** The limit of quantitation and the sample results were adjusted to reflect sample dilution.
- I1** The relative intensity of a characteristic ion in a sample analyte exceeded the acceptance with respect to the reference spectra, indicating interference.
- L1** When testing for pesticides, fungicides, herbicides, growth regulators, heavy metals, or residual solvents, the percent recovery of a laboratory control sample is greater than the acceptance limits, but the sample's target analytes were not detected above the maximum allowable concentrations for the analytes in the sample.
- M1** The recovery from the matrix spike was high, but the recovery from the laboratory control sample was within acceptance criteria.
- M2** The recovery from the matrix spike was low, but the recovery from the laboratory control sample was within acceptance criteria.
- M3** The recovery from the matrix spike was unusable because the analyte concentration was disproportionate to the spike level, but the recovery from the laboratory control sample was within acceptance criteria.
- M4** The analysis of a spiked sample required a dilution such that the spike recovery calculation does not provide useful information, but the recovery from the associated laboratory control sample was within acceptance criteria.
- M5** The analyte concentration was determined by the method of standard addition, in which the standard is added directly to the aliquots of the analyzed sample.
- M6** A description of the variance is described in the final report of testing according to R9-17- 404.06(B)(3)(d)(ii).
- Q1** Sample integrity was not maintained.
- Q2** The sample is heterogeneous, and sample homogeneity could not be readily achieved using routine laboratory practices.
- Q3** Testing result is for informational purposes only and cannot be used to satisfy dispensary testing requirements in R9-17-317.01(A) or labeling requirements in R9-17-317.
- R1** The relative percent difference for the laboratory control sample and duplicate exceeded the limit, but the recovery was within acceptance criteria.
- R2** The relative percent difference for a sample and duplicate exceeded the limit.
- V1** The recovery from continuing calibration verification standards exceeded the acceptance limits, but the sample's target analytes were not detected above the maximum allowable for the analytes in the sample.

**Notes:**

**Ahmed Munshi**

Technical Laboratory Director

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