
Test Report

Report No. : AGC12769210601-003

SAMPLE NAME : 40z Double Wall SS Vacuum Tumbler, 20oz Double Wall SS Vacuum Tumbler, 14oz double wall camping mug with SS outer, plastic liner and handle

MODEL NAME : /

APPLICANT : Promo Stock USA, LLC

STANDARD(S) : Please refer to follow page(s).

DATE OF ISSUE : Jul.07, 2021

Attestation of Global Compliance (Shenzhen) Std & Tech Co., Ltd.



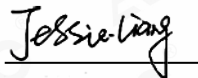
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Applicant : Promo Stock USA, LLC
Address : 418 Cloverleaf Drive, Baldwin Park, CA 91706
Test Site : 6/F., Building 2, Sanwei Chaxi Industrial Park, Sanwei Community,
Hangcheng Street, Bao'an District, Shenzhen, Guangdong, China

Report on the submitted sample(s) said to be:

Sample Name : 40z Double Wall SS Vacuum Tumbler, 20oz Double Wall SS Vacuum
Tumbler, 14oz double wall camping mug with SS outer, plastic liner and
handle
Sample Received Date : Jun.28, 2021
Testing Period : Jun.28, 2021 to Jul.07, 2021

Approved by: 

Liangdan, Jessie.Liang

Technical Director

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Test Requested:

Conclusion

- | | |
|--|---|
| <ol style="list-style-type: none"> 1. As specified by client, to determine the Lead (Pb) content in the submitted sample(s) with reference to U.S. Consumer Product Safety Improvement Act (CPSIA) of 2008 (H.R. 4040) & Amendment Act H.R. 2715, Title I, Section 101. 2. As specified by client, to determine the Phthalates content in the submitted sample(s) with reference to Section 108 of the U.S. Consumer Product Safety Improvement Act (CPSIA) of 2008(H.R. 4040) & Amendment Rule 16 CFR part 1307. 3. As specified by client, to determine the Lead Content in the submitted sample in accordance with California Proposition 65 for Lead Content Requirement. 4. As specified by client, to determine the Phthalates Content in the submitted sample in accordance with California Proposition 65 for Phthalate Content Requirement. 5. As specified by client, to determine the Lead content in the submitted sample in accordance with SOR/2018-83 (Consumer Products Containing Lead Regulations) under Canada Consumer Product Safety Act. 6. As specified by client, to determine the Bisphenol A (BPA) content in the submitted samples with reference to regulations of the United States-Tennessee, Iowa, Alaska. 7. As specified by client, to determine Chromium content in the submitted sample and in accordance with FDA GRAs 8. As specified by client, to determine the acrylonitrile extract in the submitted sample(s) with reference to FDA 21 CFR 180.22& 181.32. <ul style="list-style-type: none"> - 3% Acetic acid - 8% Ethanol - n-heptane 9. As specified by client, to determine the chloroform extract in the submitted sample(s) with reference to FDA 21 CFR 177.1210. <ul style="list-style-type: none"> - Water | <p>Pass</p> <p>Pass</p> <p>Pass</p> <p>Pass</p> <p>Pass</p> <p>Pass</p> <p>Pass</p> <p>Pass</p> <p>Pass</p> <p>Pass</p> <p>Pass</p> <p>Pass</p> |
|--|---|

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Test Result:
1. Test Result of Lead Content (for H.R. 4040)

| | |
|--------------------------------|--|
| Test Item | Lead |
| Limit(mg/kg) | ≤100(substrate), ≤90(coating or paint) |
| MDL(mg/kg) | 10 |
| Test Method/ Instrument | CPSC-CH-E1001-08.3 (metal), CPSC-CH-E1002-08.3 (nonmetal), CPSC-CH-E1003-09.1 (coating or paint) / ICP-OES |

| No. | Test result (mg/kg) | Conclusion |
|------------------|---------------------|------------|
| | Lead (Pb) | |
| 3-1 | N.D. | Conformity |
| 3-2 | N.D. | Conformity |
| 3-3 | N.D. | Conformity |
| 3-4 | N.D. | Conformity |
| 3-5 | N.D. | Conformity |
| 3-6 [△] | N.D. | Conformity |
| 3-7 [△] | N.D. | Conformity |
| 3-8 [△] | N.D. | Conformity |

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2. Test Result of Phthalate Content (for H.R. 4040)

| Test Item | Test Method/ Instrument | MDL | Limit |
|---|------------------------------|--------|-------|
| Diisobutyl phthalate(DIBP) (CAS No.: 84-69-5) | CPSC-CH-C1001-09.4/ GC-MS | 0.010% | ≤0.1% |
| Dibutyl phthalate (DBP) (CAS No.: 84-74-2) | | 0.010% | ≤0.1% |
| Butylbenzyl phthalate (BBP) (CAS No.: 85-68-7) | | 0.010% | ≤0.1% |
| Di-(2-ethylhexyl) Phthalate (DEHP) (CAS No.: 117-81-7) | | 0.010% | ≤0.1% |
| Di-isononyl phthalate (DINP) (CAS No.: 28553-12-0; 68515-48-0) | | 0.010% | ≤0.1% |
| Di-n-hexyl phthalate(DHEXP/DNHP) (CAS No.: 84-75-3) | | 0.010% | ≤0.1% |
| Di-n-pentyl phthalate(DPENP) (CAS No.:131-18-0) | | 0.010% | ≤0.1% |
| Dicyclohexly phthalate(DCHP) (CAS No.: 84-61-7) | | 0.010% | ≤0.1% |

| No. | Test result (%) | | | | | | | | Conclusion |
|------------------|-----------------|------|------|------|------|-------|-------|------|------------|
| | DIBP | DBP | BBP | DEHP | DINP | DHEXP | DPENP | DCHP | |
| 3-4 | N.D. | N.D. | N.D. | N.D. | N.D. | N.D. | N.D. | N.D. | Conformity |
| 3-5 | N.D. | N.D. | N.D. | N.D. | N.D. | N.D. | N.D. | N.D. | Conformity |
| 3-6 ^Δ | N.D. | N.D. | N.D. | N.D. | N.D. | N.D. | N.D. | N.D. | Conformity |
| 3-7 ^Δ | N.D. | N.D. | N.D. | N.D. | N.D. | N.D. | N.D. | N.D. | Conformity |
| 3-8 ^Δ | N.D. | N.D. | N.D. | N.D. | N.D. | N.D. | N.D. | N.D. | Conformity |

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3. Test Result of Lead Content (for California Proposition 65)

| | |
|---|---|
| Test Item | Lead |
| Limit (Client's Requirement) (mg/kg) | 100(substrate), 90(coating or paint) |
| MDL(mg/kg) | 10 |
| Test Method/ Instrument | CPSC-CH-E1001-08.3 (metal), CPSC-CH-E1002-08.3 (nonmetal), CPSC-CH-E1003-09.1 (coating) / ICP-OES |

| No. | Test result (mg/kg) | Conclusion |
|------------------|---------------------|------------|
| | Lead (Pb) | |
| 3-1 | N.D. | Conformity |
| 3-2 | N.D. | Conformity |
| 3-3 | N.D. | Conformity |
| 3-4 | N.D. | Conformity |
| 3-5 | N.D. | Conformity |
| 3-6 [△] | N.D. | Conformity |
| 3-7 [△] | N.D. | Conformity |
| 3-8 [△] | N.D. | Conformity |

4. Test Result of Phthalate Content (for California Proposition 65)

| Test Item | Test Method/ Instrument | MDL | Limit (Client's Requirement) |
|---|------------------------------|--------|------------------------------|
| Dibutyl phthalate (DBP) (CAS No.: 84-74-2) | CPSC-CH-C1001-09.4/ GC-MS | 0.010% | 0.1% |
| Butylbenzyl phthalate (BBP) (CAS No.: 85-68-7) | | 0.010% | 0.1% |
| Di-(2-ethylhexyl) Phthalate (DEHP) (CAS No.: 117-81-7) | | 0.010% | 0.1% |
| Di-n-octyl phthalate (DNOP) * (CAS No.: 117-84-0) | | 0.010% | 0.1% |
| Di-isononyl phthalate (DINP) (CAS No.: 28553-12-0; 68515-48-0) | | 0.010% | 0.1% |
| Di-isodecyl phthalate(DIDP) (CAS No.: 26761-40-0; 68515-49-1) | | 0.010% | 0.1% |
| Di-n-hexyl phthalate(DNHP) (CAS No.: 84-75-3) | | 0.010% | 0.1% |

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| No. | Test result (%) | | | | | | | Conclusion |
|------------------|-----------------|------|------|------|------|------|------|------------|
| | DBP | BBP | DEHP | DNOP | DINP | DIDP | DNHP | |
| 3-3 | N.D. | N.D. | N.D. | N.D. | N.D. | N.D. | N.D. | Conformity |
| 3-4 | N.D. | N.D. | N.D. | N.D. | N.D. | N.D. | N.D. | Conformity |
| 3-5 | N.D. | N.D. | N.D. | N.D. | N.D. | N.D. | N.D. | Conformity |
| 3-6 [△] | N.D. | N.D. | N.D. | N.D. | N.D. | N.D. | N.D. | Conformity |
| 3-7 [△] | N.D. | N.D. | N.D. | N.D. | N.D. | N.D. | N.D. | Conformity |
| 3-8 [△] | N.D. | N.D. | N.D. | N.D. | N.D. | N.D. | N.D. | Conformity |

5. Test Result of Lead Content (for SOR/2018-83)

| | |
|-------------------------|------------------------------------|
| Test Item | Lead(Pb) |
| Limit(mg/kg) | ≤90 |
| MDL(mg/kg) | 5 |
| Test Method/ Instrument | Health Canada Method C-02/ ICP-OES |

| No. | Test result (mg/kg) | Conclusion |
|------------------|---------------------|------------|
| | Lead(Pb) | |
| 3-1 | N.D. | Conformity |
| 3-2 | N.D. | Conformity |
| 3-3 | N.D. | Conformity |
| 3-4 | N.D. | Conformity |
| 3-5 | N.D. | Conformity |
| 3-6 [△] | N.D. | Conformity |
| 3-7 [△] | N.D. | Conformity |
| 3-8 [△] | N.D. | Conformity |

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6. Test Result of Bisphenol A (BPA) Content (United States)

| | |
|--------------------------------|---|
| Test Item | Bisphenol A (BPA) |
| Limit (mg/kg) | Absent |
| MDL(mg/kg) | 1 |
| Test Method/ Instrument | EPA 3540C:1996 & EPA 8321B:2007/ LC-MS-MS |

| No. | Test result (mg/kg) | Conclusion |
|-----|---------------------|------------|
| | Bisphenol A (BPA) | |
| 3-3 | N.D. | Conformity |
| 3-4 | N.D. | Conformity |

7. Test Result of Chromium (Cr) Content

| Test Items | Test Condition | Unit | MDL | Result | Limit |
|-------------------|---|------|-------|-------------------|-------|
| | | | | 3-2 | |
| Chromium (Cr) | EPA 3050B:1996&EPA 6010D:2018/ ICP-OES | % | 0.001 | 16.395 | ≥16 |
| Conclusion | / | / | / | Conformity | / |

8. Test result of FDA 21 CFR 180.22& 181.32

| Test Items | Test Condition | Unit | MDL | Result | Limit |
|-----------------------|---|--------------------|-------|-------------------|-------|
| | | | | 3-3 | |
| acrylonitrile extract | 3% Acetic acid, Boil and cool to 37.8 °C | mg/in ² | 0.001 | N.D. | 0.003 |
| | 8% Ethanol, 150°F, 2h | | 0.001 | N.D. | 0.003 |
| | n-heptane, 180°F, 15min | | 0.001 | N.D. | 0.003 |
| Conclusion | / | / | / | Conformity | / |

9. Test Result of FDA 21 CFR 177.1210

| Test Item(s) | Test Condition | Unit | MDL | Result(s) | Limit |
|--------------------|--------------------------|-------|-----|-------------------|-------|
| | | | | 3-4 | |
| chloroform extract | Boil and cool to 37.8 °C | mg/kg | 5 | N.D. | 50 |
| Conclusion | / | / | / | Conformity | / |

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Note:

% = percentage

mg/kg = milligram per kilogram

N.D.= Not Detected (lower than method detection limit)

mg/in²=Milligrams per square inch

MDL = Method detection limit

Remark:

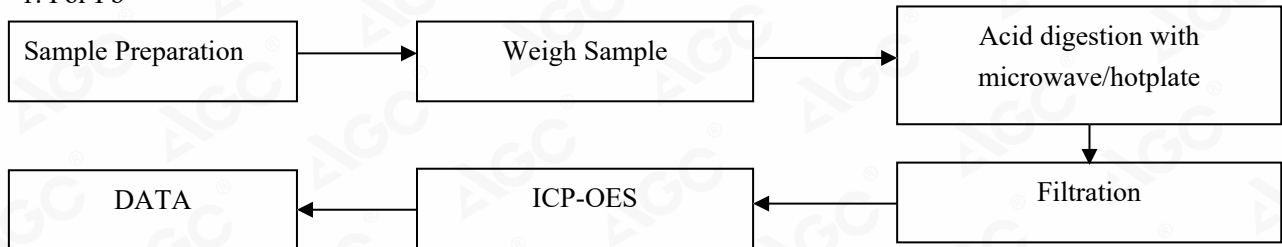
- *= As specified by client, this item is extra testing, this item is not included in above test requested.
- Δ= As specified by client, the submitted samples were mixed to test.

Sample Description:

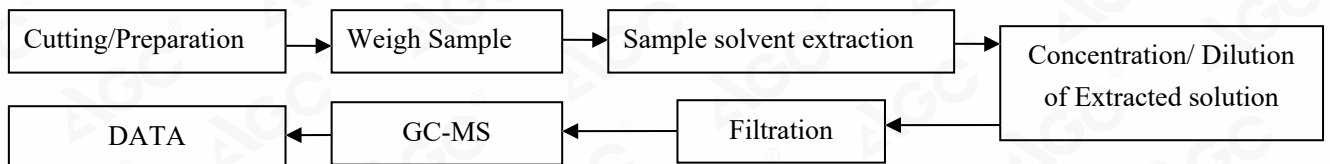
| | |
|-----|---|
| 3-1 | Outer 201 stainless steel |
| 3-2 | Inner 304 stainless steel |
| 3-3 | Transparent as cover |
| 3-4 | Silicone ring (The sample volume was 500 mL) |
| 3-5 | Black PP handle |
| 3-6 | Blue coating + green coating |
| 3-7 | Grey coating + black coating |
| 3-8 | White coating + Red Coating + dark blue coating |

Test Flow Chart

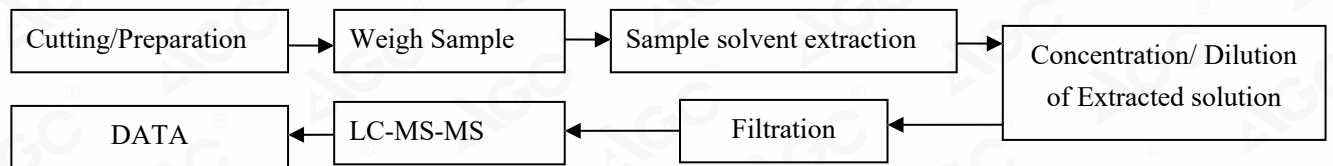
1. For Pb



2. For Phthalate



3. For Bisphenol A (BPA)



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The photo of the sample



AGC12769210601-003

AGC authenticate the photo only on original report

*** End of Report ***

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Conditions of Issuance of Test Reports

1. All samples and goods are accepted by the Attestation of Global Compliance (Shenzhen) Std & Tech Co., Ltd. (the “Company”) solely for testing and reporting in accordance with the following terms and conditions. The company provides its services on the basis that such terms and conditions constitute express agreement between the company and any person, firm or company requesting its services (the “Clients”).
2. Any report issued by Company as a result of this application for testing services (the “Report”) shall be issued in confidence to the Clients and the Report will be strictly treated as such by the Company. It may not be reproduced either in its entirety or in part and it may not be used for advertising or other unauthorized purposes without the written consent of the Company. The Clients to whom the Report is issued may, however, show or send it, or a certified copy thereof prepared by the Company to its customer, supplier or other persons directly concerned. The Company will not, without the consent of the Clients, enter into any discussion or correspondence with any third party concerning the contents of the Report, unless required by the relevant governmental authorities, laws or court orders.
3. The Company shall not be called or be liable to be called to give evidence or testimony on the Report in a court of law without its prior written consent, unless required by the relevant governmental authorities, laws or court orders.
4. The non-CMA report issued by AGC is only permitted to be used by the client as internal reference use and shall not be used for public demonstration purpose.
5. In the event of the improper use of the report as determined by the Company, the Company reserves the right to withdraw it, and to adopt any other additional remedies which may be appropriate.
6. Samples submitted for testing are accepted on the understanding that the Report issued cannot form the basis of, or be the instrument for, any legal action against the Company.
7. The Company will not be liable for or accept responsibility for any loss or damage however arising from the use of information contained in any of its Reports or in any communication whatsoever about its said tests or investigations.
8. Clients wishing to use the Report in court proceedings or arbitration shall inform the Company to that effect prior to submitting the sample for testing.
9. The Company is not responsible for recalling the electronic version of the original report when any revision is made to them. The Client assumes the responsibility to providing the revised version to any interested party who uses them.
10. Subject to the variable length of retention time for test data and report stored hereinto as otherwise specifically required by individual accreditation authorities, the Company will only keep the supporting test data and information of the test report for a period of six years. The data and information will be disposed of after the aforementioned retention period has elapsed. Under no circumstances shall we provide any data and information which has been disposed of after retention period. Under no circumstances shall we be liable for damage of any kind, including (but not limited to) compensatory damages, lost profits, lost data, or any form of special, incidental, indirect, consequential or punitive damages of any kind, whether based on breach of contract of warranty, tort (including negligence), product liability or otherwise, even if we are informed in advance of the possibility of such damages.

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