Test Report

Report No.	:	AGC12769210601-002

SAMPLE NAME	:	17oz Double Wall SS Vacuum Bottle
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	:	17oz Double Wall SS Vacuum Bottle
Sample Received Date	:	Jun.28, 2021
Testing Period	:	Jun.28, 2021 to Jul.07, 2021

Approved by: Jessie.Liang

Liangdan, Jessie.Liang

Technical Director

Test Requested:	Conclusion
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.	**Pass**
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.	**Pass**
3. As specified by client, to determine the Lead Content in the submitted sample in accordance with California Proposition 65 for Lead Content Requirement.	**Pass**
4. As specified by client, to determine the Phthalates Content in the submitted sample in accordance with California Proposition 65 for Phthalate Content Requirement.	**Pass**
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.	**Pass**
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.	**Pass**
7. As specified by client, to determine Chromium content in the submitted sample and in accordance with FDA GRAs	**Pass**
8. As specified by client, the following items are determined in the submitted sample with reference to the US Food And Drug Administration (FDA) the food contact test standard: 21 CFR 177.1520:	
- N-hexane extractives	**Pass**
- Xylene extractives	**Pass**
- Density	**Pass**
- Melting point	**Pass**
9. As specified by client, to determine the chloroform extract in the submitted sample(s) with reference to FDA 21 CFR 177.1210.	
- Water	**Pass**

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)	
2-1	N.D.	**Conformity**
2-2	N.D.	**Conformity**
2-3	N.D.	**Conformity**
2-4	N.D.	**Conformity**
2-5△	N.D.	**Conformity**

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	
2-3	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	**Conformity**
2-4	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	**Conformity**
2-5△	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	**Conformity**

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)	
2-1	N.D.	**Conformity**
2-2	N.D.	**Conformity**
2-3	N.D.	**Conformity**
2-4	N.D.	**Conformity**
2-5△	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%

No.	**Test result (%)**							**Conclusion**
	DBP	**BBP**	**DEHP**	**DNOP**	**DINP**	**DIDP**	**DNHP**	
2-3	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	**Conformity**
2-4	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	**Conformity**
2-5△	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)	
2-1	N.D.	**Conformity**
2-2	N.D.	**Conformity**
2-3	N.D.	**Conformity**
2-4	N.D.	**Conformity**
2-5△	N.D.	**Conformity**

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)	
2-3	N.D.	**Conformity**
2-4	N.D.	**Conformity**

7. Test Result of Chromium (Cr) Content

Test Items	**Test Condition**	Unit	**MDL**	**Result**	**Limit**
				2-2	
Chromium (Cr)	EPA 3050B:1996&EPA 6010D:2018/ ICP-OES	%	0.001	18.489	≥16
Conclusion	/	/	/	**Conformity**	/

8. Test Result of FDA 21 CFR 177.1520

Test Item	Test condition	Unit	MDL	Result(s)	Limit
				2-3	
N-hexane extractives	Reflux, 2h	%	0.1	0.2	6.4
Xylene extractives	Reflux, 2h	%	0.1	1.2	9.8
Density	/	g/cm^3	/	0.906	0.880~0.913
Melting point	/	℃	/	169.7	160~180
Conclusion	/	/	/	**Conformity**	/

9. Test Result of FDA 21 CFR 177.1210

Test Item(s)	Test Condition	Unit	MDL	Result(s)	Limit
				2-4	
chloroform extract	Boil and cool to 37.8 ℃	mg/kg	5	N.D.	50
Conclusion	/		/	**Conformity**	/

Note:

% = percentage

g/cm^3 = grams per cubic centimeter

mg/kg = milligram per kilogram

MDL = Method detection limit

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

Remark:

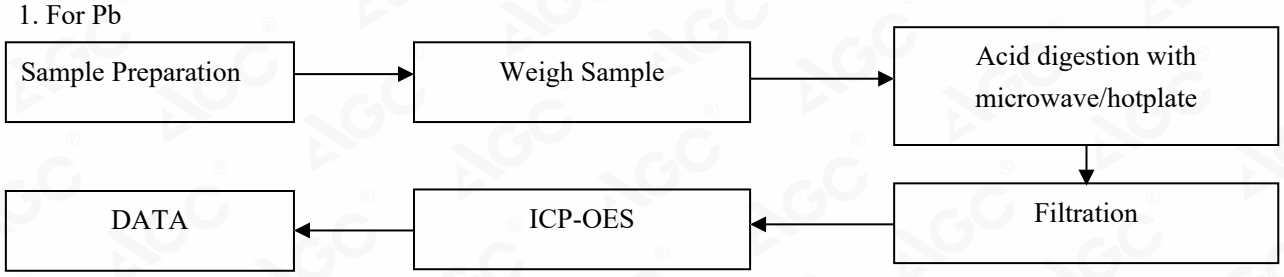
- *= As specified by client, this item is extra testing, this item is not included in above test requested.
- △= As specified by client, the submitted samples were mixed to test.

Sample Description:

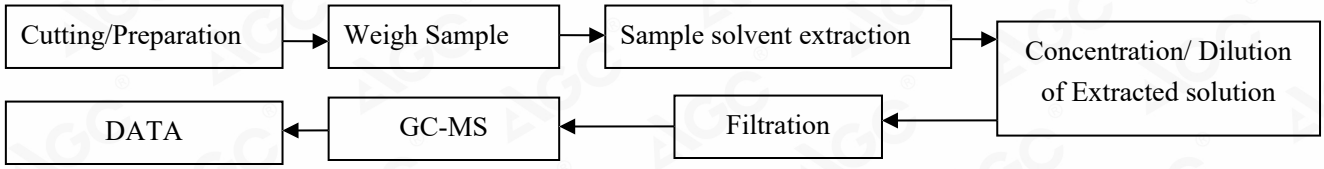
2-1	Outer 201 stainless steel
2-2	Inner 304 stainless steel
2-3	Black PP cover
2-4	Silicone ring (The sample volume was 530 mL)
2-5	Blue coating + black coating + white coating

Test Flow Chart

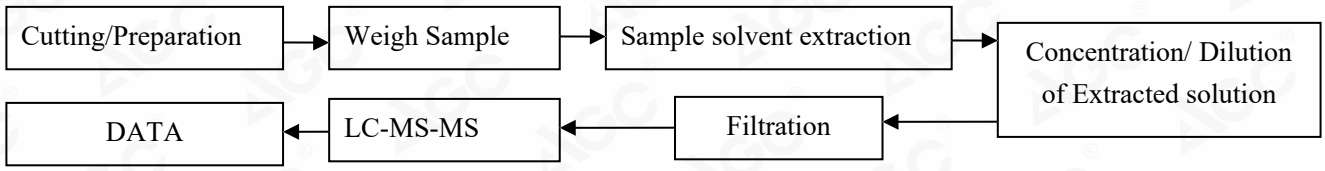
1. For Pb



2. For Phthalate



3. For Bisphenol A (BPA)



The photo of the sample

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AGC authenticate the photo only on original report

*** End of Report ***

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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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