



TEST REPORT

Reference No. : SZ2025040945-2E

Date : Apr. 28, 2025

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Client : Mahlwerck Porzellan GmbH

Address : An der Alten Spinnerei 1 D-83059 Kolbermoor

The following merchandise was (were) submitted and identified by the client as:

Name of Product : Audi Cups 3 colors

Test Model : shape 147

Model May Cover : /

Main Material: Porcelain

Lot No.: 04-2025

Buyer: Cyberwear GmbH

Sample Received : Apr. 21, 2025

Test Period : Apr. 21, 2025 - Apr. 28, 2025

Test Specification and Conclusion:

Please refer to next page

Prepared By :

Seth Kong
Testing Engineer

Reviewed By :

Dora Cheng
Reporter Supervisor

Issued By :



Ada Wang
Lab Manager



SZ2025040945-2E

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Test Request		Conclusion
1.	Leachable Lead, Cadmium content according to California Proposition 65 for Ceramicware - Food and Beverage use products (Internal), Consent judgement No. SF -938430	PASS
2.	Leachable Lead, Cadmium content according to California Proposition 65 for food and beverage use products (lip and rim area), Consent judgement No. SF-436429 & SF-440811	PASS
3.	Lead and Cadmium content in surface wipe sample according to California Proposition 65 for ceramic ware - food and beverage use products, Consent judgement no. SF 436429/SF- 440811	PASS
4.	Extractable Lead and Cadmium content according to FDA 7117.07&7117.06	PASS
5.	Extractable Lead, Cadmium content for Ceramic according to Japan Ministry of Health and Welfare Ordinance No. 370, Specifications and Standards for Foods, Food Additives, etc. Under The Food Sanitation Law	PASS
6.	Extractable Lead and Cadmium content according to SOR/2016-175	PASS
7.	Cadmium (Cd) content according to EC Regulation 1907/2006, REACH Annex XVII (entry 23)	PASS
8.	Lead (Pb) content according to EC Regulation 1907/2006, REACH Annex XVII (entry 63)	PASS
9.	Lead(Pb) content according to Canada Consumer Products Containing Lead (Contact with Mouth) Regulations: SOR/2018-83	PASS
10.	Polycyclic Aromatic Hydrocarbons (PAHs) contents according to AfPS GS 2019:01 PAK	PASS
11.	Polycyclic Aromatic Hydrocarbons (PAHs) contents according to EC Regulation 1907/2006, (EU) No 1272/2013 Amending PAHs of REACH Annex XVII(entry 50)	PASS
12.	Phthalate content according to EC Regulation 1907/2006, REACH Annex XVII (entry 51&52)	PASS
13.	Bisphenol-A (BPA) content according to US California Proposition 65	PASS
14.	Phthalates content according to US California Proposition 65& California Bill AB 1108:2007& Consumer Product Safety Improvement Act of 2008, CPSIA section 108	PASS
European Council Directive 84/500/EEC and its amendment 2005/31/EC - Amending Council Directive 84/500/EEC as regards a declaration of compliance and performance criteria of the analytical method for ceramic articles intended to come into contact with foodstuffs, General Requirement (Article 3) in EU Regulation No. 1935/2004		
15.	Extractable Lead, Cadmium and Cobalt content for Ceramic	PASS
German Food, Articles of Daily Use and Feed Code of September 1, 2005(LFGB), Section 30 and DIN		

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51032:2017-07, ALS Opinion Number 2017/15, General Requirement (Article 3) in EU Regulation No.1935/2004

16.	Sensorial examination odour and taste test for full article	PASS
17.	Extractable Lead, Cadmium and Cobalt content for Ceramic	PASS

***** To be continued *****



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TEST RESULTS:

1. Leachable Lead, Cadmium content

Test Method: With reference to AOAC 16th Ed. (2000) Section 973.32, Analysis was performed by Atomic Absorption Spectrometer (AAS).

No.1#-1	Depth (mm)	Diameter (mm)	Volume of Leaching Solution (ml)	Lead (mg/L)	Cadmium (mg/L)
1	86	79	380	<0.05	<0.01
2	86	79	380	<0.05	<0.01
3	86	79	380	<0.05	<0.01
4	86	79	380	<0.05	<0.01
5	86	79	380	<0.05	<0.01
6	86	79	380	<0.05	<0.01
California Proposition 65:				0.100	0.189

No.2#-1	Depth (mm)	Diameter (mm)	Volume of Leaching Solution (ml)	Lead (mg/L)	Cadmium (mg/L)
1	85	78	380	<0.05	<0.01
2	85	78	380	<0.05	<0.01
3	85	78	380	<0.05	<0.01
4	85	78	380	<0.05	<0.01
5	85	78	380	<0.05	<0.01
6	85	78	380	<0.05	<0.01
California Proposition 65:				0.100	0.189

No.3#-1	Depth (mm)	Diameter (mm)	Volume of Leaching Solution (ml)	Lead (mg/L)	Cadmium (mg/L)
1	85	78	380	<0.05	<0.01
2	85	78	380	<0.05	<0.01
3	85	78	380	<0.05	<0.01
4	85	78	380	<0.05	<0.01
5	85	78	380	<0.05	<0.01
6	85	78	380	<0.05	<0.01
California Proposition 65:				0.100	0.189

Remark: The above California limits were quoted from the consent judgement No. SF-938430 settled by superior court of the state of California for the city and county of San Francisco, For ceramicware STQ Testing Services Co., Ltd.

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Food and Beverage use products (Internal) based on the California Proposition 65.

2. Leachable Lead, Cadmium content

Test Method: With reference to ASTM C927-80(2019)e1, Standard Methods of Test for Lead and Cadmium extracted from Glazed Ceramic Surface of the Lip and Rim Area, Analysis was performed by Atomic Absorption Spectrometer (AAS).

No.1#-2	Depth (mm)	Diameter (mm)	Volume of Leaching Solution (ml)	Lead (mg/kg)	Cadmium (mg/kg)
1	86	79	270	<0.05	<0.01
2	86	79	270	<0.05	<0.01
3	86	79	270	<0.05	<0.01
4	86	79	270	<0.05	<0.01
5	86	79	270	<0.05	<0.01
6	86	79	270	<0.05	<0.01
California Proposition 65:				0.5	4.0

No.2#-2	Depth (mm)	Diameter (mm)	Volume of Leaching Solution (ml)	Lead (mg/kg)	Cadmium (mg/kg)
1	85	78	270	<0.05	<0.01
2	85	78	270	<0.05	<0.01
3	85	78	270	<0.05	<0.01
4	85	78	270	<0.05	<0.01
5	85	78	270	<0.05	<0.01
6	85	78	270	<0.05	<0.01
California Proposition 65:				0.5	4.0

No.3#-2	Depth (mm)	Diameter (mm)	Volume of Leaching Solution (ml)	Lead (mg/kg)	Cadmium (mg/kg)
1	85	78	270	<0.05	<0.01
2	85	78	270	<0.05	<0.01
3	85	78	270	<0.05	<0.01
4	85	78	270	<0.05	<0.01
5	85	78	270	<0.05	<0.01
6	85	78	270	<0.05	<0.01
California Proposition 65:				0.5	4.0

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Remark: The above limits were quoted from the consent judgement No. SF 436429 & SF-440811 settled by superior court of the state of California for the city and county of San Francisco, for food and beverage use products (lip and rim area) based on the California Proposition 65.

3. Lead and Cadmium Content

Test Method: With reference to NIOSH 9100, Analysis was performed by GF-AAS.

Test Item(s)	Unit	MDL	Test Results	Limited Value***
			1#-1	
Lead (Pb)	µg/article	0.1	N.D.	1
Cadmium (Cd)	µg/article	0.1	N.D.	4

Test Item(s)	Unit	MDL	Test Results	Limited Value***
			2#-1	
Lead (Pb)	µg/article	0.1	N.D.	1
Cadmium (Cd)	µg/article	0.1	N.D.	4

Test Item(s)	Unit	MDL	Test Results	Limited Value***
			3#-1	
Lead (Pb)	µg/article	0.1	N.D.	1
Cadmium (Cd)	µg/article	0.1	N.D.	4

Remark: ***The limit was quoted from the consent judgement no. SF- 436429/5F 440811 settled by superior court of the state of California for the city and county of San Francisco, for ceramicware – food and beverage use products based on the California Proposition 65.

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4. Extractable Lead and Cadmium Content

Test Method: With reference to AOAC 16th Ed. (2000) Section 973.32, Analysis was performed by Atomic Absorption Spectrometer (AAS).

No. 1#-1	Depth (mm)	Diameter (mm)	Volume of Leaching Solution (ml)	Lead (µg/ml)	Cadmium (µg/ml)
1	86	79	380	< 0.1	< 0.01
2	86	79	380	< 0.1	< 0.01
3	86	79	380	< 0.1	< 0.01
4	86	79	380	< 0.1	< 0.01
5	86	79	380	< 0.1	< 0.01
6	86	79	380	< 0.1	< 0.01
Any one of six of FDA limit				0.5	0.5

No. 2#-1	Depth (mm)	Diameter (mm)	Volume of Leaching Solution (ml)	Lead (µg/ml)	Cadmium (µg/ml)
1	85	78	380	< 0.1	< 0.01
2	85	78	380	< 0.1	< 0.01
3	85	78	380	< 0.1	< 0.01
4	85	78	380	< 0.1	< 0.01
5	85	78	380	< 0.1	< 0.01
6	85	78	380	< 0.1	< 0.01
Any one of six of FDA limit				0.5	0.5

No. 3#-1	Depth (mm)	Diameter (mm)	Volume of Leaching Solution (ml)	Lead (µg/ml)	Cadmium (µg/ml)
1	85	78	380	< 0.1	< 0.01
2	85	78	380	< 0.1	< 0.01
3	85	78	380	< 0.1	< 0.01
4	85	78	380	< 0.1	< 0.01
5	85	78	380	< 0.1	< 0.01
6	85	78	380	< 0.1	< 0.01
Any one of six of FDA limit				0.5	0.5

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Permissible Limit

Category	Criteria	Unit	Permissible limit
			Lead
Flatware	average of 6 units	µg/ml	3.0
Small Hollowware other than cups and mugs	any one of 6 units	µg/ml	2.0
Cups/mugs	any one of 6 units	µg/ml	0.5
Large Hollowware other than pitchers	any one of 6 units	µg/ml	1.0
Pitchers	any one of 6 units	µg/ml	0.5

Category	Criteria	Unit	Permissible limit
			Cadmium
Flatware	average of 6 units	µg/ml	0.5
Small Hollowware	any one of 6 units	µg/ml	0.5
Large Hollowware	any one of 6 units	µg/ml	0.25

Remark:

Flatware: ceramic articles which have an internal depth as measured from the lowest point to the horizontal plane passing through the upper rim, that does not exceed 25 mm.

Hollowware: ceramic articles which have an internal depth measured from the lowest point to the horizontal plane passing through the upper rim, greater than 25 mm.

Small hollowware: a capacity of less than 1.1 liter.

Large hollowware: a capacity of 1.1 liter or more.

5. Extractable Lead, Cadmium Content

Test Method: Followed by analysis using Inductively Coupled Argon Plasma-Atomic Emission Spectrometer.

Sample No. : 1#-1	Result				Limit
	1	2	3	4	
Released Lead (mg/l)	< 0.1	< 0.1	< 0.1	< 0.1	2
Released Cadmium (mg/l)	< 0.01	< 0.01	< 0.01	< 0.01	0.5
Volume of 4% Acetic acid used (ml)	380	380	380	380	---
Depth (mm)	86	86	86	86	---
Diameter (mm)	79	79	79	79	---

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Sample No. : 2#-1	Result				Limit
	1	2	3	4	
Released Lead (mg/l)	< 0.1	< 0.1	< 0.1	< 0.1	2
Released Cadmium (mg/l)	< 0.01	< 0.01	< 0.01	< 0.01	0.5
Volume of 4% Acetic acid used (ml)	380	380	380	380	---
Depth (mm)	85	85	85	85	---
Diameter (mm)	78	78	78	78	---

Sample No. : 3#-1	Result				Limit
	1	2	3	4	
Released Lead (mg/l)	< 0.1	< 0.1	< 0.1	< 0.1	2
Released Cadmium (mg/l)	< 0.01	< 0.01	< 0.01	< 0.01	0.5
Volume of 4% Acetic acid used (ml)	380	380	380	380	---
Depth (mm)	85	85	85	85	---
Diameter (mm)	78	78	78	78	---

Ceramic	Samples less than 2.5cm deep after liquid is filled or those not capable to be filled in.			Cadmium Lead	not more than 0.7 µg/cm ² not more than 8 µg/cm ²
	Samples deeper than 2.5cm when filled	Implements other than for cooking by heating	Capacity less than 1.1 L	Cadmium Lead	not more than 0.5 µg/ml not more than 2 µg/ml
			Cap. between 1L and 3L	Cadmium Lead	not more than 0.25 µg/ml not more than 1 µg/ml
			Cap. not less than 3L	Cadmium Lead	not more than 0.25 µg/ml not more than 0.5 µg/ml
		Implements used for cooking by heating		Cadmium Lead	not more than 0.05 µg/ml not more than 0.5 µg/ml

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6. Extractable Lead and Cadmium Content

Test Method: With reference to ISO 6486-1:2019, Analysis was performed by Atomic Absorption Spectrometer (AAS).

No. 1#-1	Depth (mm)	Diameter (mm)	Volume of Leaching Solution (ml)	Lead (mg/L)	Cadmium (mg/L)
1	86	79	380	< 0.1	< 0.01
2	86	79	380	< 0.1	< 0.01
3	86	79	380	< 0.1	< 0.01
4	86	79	380	< 0.1	< 0.01
5	86	79	380	< 0.1	< 0.01
6	86	79	380	< 0.1	< 0.01
SOR/2016-175 limit				0.5	0.5

No. 2#-1	Depth (mm)	Diameter (mm)	Volume of Leaching Solution (ml)	Lead (mg/L)	Cadmium (mg/L)
1	85	78	380	< 0.1	< 0.01
2	85	78	380	< 0.1	< 0.01
3	85	78	380	< 0.1	< 0.01
4	85	78	380	< 0.1	< 0.01
5	85	78	380	< 0.1	< 0.01
6	85	78	380	< 0.1	< 0.01
SOR/2016-175 limit				0.5	0.5

No. 3#-1	Depth (mm)	Diameter (mm)	Volume of Leaching Solution (ml)	Lead (mg/L)	Cadmium (mg/L)
1	85	78	380	< 0.1	< 0.01
2	85	78	380	< 0.1	< 0.01
3	85	78	380	< 0.1	< 0.01
4	85	78	380	< 0.1	< 0.01
5	85	78	380	< 0.1	< 0.01
6	85	78	380	< 0.1	< 0.01
SOR/2016-175 limit				0.5	0.5

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Permissible Limit

Category	Unit	Permissible limit	
		Lead	Cadmium
Flatware	mg/L	3.0	0.5
Small Hollowware other than cups and mugs	mg/L	2.0	0.5
Cups/mugs	mg/L	0.5	0.5
Large Hollowware other than pitchers	mg/L	1.0	0.25
Pitchers	mg/L	0.5	0.25

7. Cadmium (Cd) Content

Test Method: With reference to EN1122:2001, Analysis was performed by ICP-AES.

Test Item(s)	MDL (mg/kg)	Test Results (mg/kg)			Limited Value* (mg/kg)
		4#	5#	6#	
Cadmium (Cd)	10	N.D.	N.D.	N.D.	1000

Remark: *The Limited value is based on EC Regulation 1907/2006, REACH Annex XVII (entry 23).

8. Lead(Pb) Content

Test Method: With reference to EPA 3052-1996 & EPA 6010D-2018, Analysis was performed by ICP-AES.

Test Item	MDL (mg/kg)	Test Results (mg/kg)			Limited Value** (mg/kg)
		4#	5#	6#	
Lead(Pb)	10	N.D.	N.D.	N.D.	500

Remark: **The Limited value is based on EC Regulation 1907/2006, REACH Annex XVII (entry 63).

9. Total Lead Content

Test Method: With reference to Canada method C-02.2.2: 2020, Analysis was performed by ICP-AES.

Test Item	MDL (mg/kg)	Test Results (mg/kg)			Limited Value*** (mg/kg)
		4#	5#	6#	
Total Lead	10	N.D.	N.D.	N.D.	90

Remark: ***The Limited value is based on Canada Consumer Products Containing Lead (Contact with Mouth) Regulations: SOR/2018-83.

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10.PAHs Contents

Test Method: With reference to AfPS GS 2019:01 PAK, Analysis was performed by GC-MS.

Test Items	Unit	MDL	Test Results		
			4#	5#	6#
Naphthalene	mg/kg	0.1	0.2	0.1	0.2
Phenanthrene		0.1	0.2	0.4	0.2
Anthracene		0.1	N.D.	N.D.	N.D.
Fluoranthene		0.1	N.D.	N.D.	N.D.
Pyrene		0.1	N.D.	N.D.	N.D.
Benzo[a]anthracene		0.1	N.D.	N.D.	N.D.
Chrysene		0.1	N.D.	N.D.	N.D.
Benzo[b]fluoranthene		0.1	N.D.	N.D.	N.D.
Benzo[k]fluoranthene		0.1	N.D.	N.D.	N.D.
Benzo[a]pyrene		0.1	N.D.	N.D.	N.D.
Indeno[1,2,3-cd]pyrene		0.1	N.D.	N.D.	N.D.
Dibenzo[a,h]anthracene		0.1	N.D.	N.D.	N.D.
Benzo[g,h,i]perylene		0.1	N.D.	N.D.	N.D.
Benzo[j]fluoranthene		0.1	N.D.	N.D.	N.D.
Benzo[e]pyrene		0.1	N.D.	N.D.	N.D.
Sum of 15 PAHs		---	0.4	0.5	0.4

***** To be continued *****

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LIMITS(mg/kg) FOR PAH IN PRODUCTS:

Parameter	Category 1	Category2		Category 3	
	Materials intended to be placed in the mouth, or materials coming into long-term contact with skin (more than 30s) during the intended use - in toys according to Directive 2009/48/EC or - for the use by children up to 3 years of age	Materials not covered by category 1, coming into long-term contact (more than 30s) or short-term repetitive contact with skin during the intended or foreseeable use		Materials covered neither by category 1 nor by category 2, coming into short-term contact (up to 30s) with skin during the intended or foreseeable use	
		a. use by children	b. other consumer products	a. use by children	b. other consumer products
Benzo[a]pyrene	< 0.2	< 0.2	< 0.5	< 0.5	< 1
Benzo[e]pyrene	< 0.2	< 0.2	< 0.5	< 0.5	< 1
Benzo[a]anthracene	< 0.2	< 0.2	< 0.5	< 0.5	< 1
Benzo[b]fluoranthene	< 0.2	< 0.2	< 0.5	< 0.5	< 1
Benzo[j]fluoranthene	< 0.2	< 0.2	< 0.5	< 0.5	< 1
Benzo[k]fluoranthene	< 0.2	< 0.2	< 0.5	< 0.5	< 1
Chrysene	< 0.2	< 0.2	< 0.5	< 0.5	< 1
Dibenzo[a,h]anthracene	< 0.2	< 0.2	< 0.5	< 0.5	< 1
Benzo[g,h,i]perylene	< 0.2	< 0.2	< 0.5	< 0.5	< 1
Indeno[1,2,3-cd]pyrene	< 0.2	< 0.2	< 0.5	< 0.5	< 1
Phenanthrene, Pyrene, Anthracene, Fluoranthene	Total < 1	Total < 5	Total < 10	Total < 20	Total < 50
Naphthalene	< 1	< 2		< 10	
Sum of 15 PAHs	< 1	< 5	< 10	< 20	< 50

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11.PAHs Contents

Test Method: With reference to AfPS GS 2019:01 PAK, Analysis was performed by GC-MS.

Test Items	CAS No.	MDL (mg/kg)	Test Results (mg/kg)			Limited Value**** (mg/kg)	
			4#	5#	6#	Category I	Category II
Benzo[a]pyrene	50-32-8	0.1	N.D.	N.D.	N.D.	1	0.5
Benzo[e]pyrene	192-97-2	0.1	N.D.	N.D.	N.D.	1	0.5
Benzo[a]anthracene	56-55-3	0.1	N.D.	N.D.	N.D.	1	0.5
Chrysene	218-01-9	0.1	N.D.	N.D.	N.D.	1	0.5
Benzo[b]fluoranthene	205-99-2	0.1	N.D.	N.D.	N.D.	1	0.5
Benzo[j]fluoranthene	205-82-3	0.1	N.D.	N.D.	N.D.	1	0.5
Benzo[k]fluoranthene	207-08-9	0.1	N.D.	N.D.	N.D.	1	0.5
Dibenzo[a,h]anthracene	53-70-3	0.1	N.D.	N.D.	N.D.	1	0.5

Remark: ****The Limited value is based on (EU) No 1272/2013 Amending PAHS of REACH Annex XVII(entry 50).

LIMITS FOR PAH :

Parameter	Category I	Category II
PAHs	Such articles include amongst other: 1. Sport equipment such as bicycles, golf clubs, racquets 2. House-hold utensils, trolleys, walking frames 3. tools for domestic use 4. clothing, footwear, gloves and sportwear 5. watch-straps, wrist-bands, masks, head-bands	Toys, including activity toys, and childcare articles

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12. Phthalates Content

Test Method: With reference to EN 14372-2004, Analysis was performed by GC-MS.

(12-1) For plasticized materials (including toys and childcare articles)

Test Item(s)	Unit	MDL	Test Results			Limited Value*****
			4#	5#	6#	
Dibutyl Phthalate(DBP)	%	0.005	N.D.	N.D.	N.D.	---
Benzylbutyl Phthalate(BBP)		0.005	N.D.	N.D.	N.D.	---
Di-(2-ethylhexyl)Phthalate(DEHP)		0.005	N.D.	N.D.	N.D.	---
Di-isobutyl phthalate (DIBP)		0.005	N.D.	N.D.	N.D.	---
Total (DBP+BBP+DEHP+DIBP)		---	< 0.020	< 0.020	< 0.020	< 0.1

(12-2) For toys and childcare articles that can be mouthed

Test Item(s)	Unit	MDL	Test Results			Limited Value*****
			4#	5#	6#	
Di-iso-nonylphthalate(DINP)	%	0.005	N.D.	N.D.	N.D.	---
Di-n-octylphthalate(DNOP)		0.005	N.D.	N.D.	N.D.	---
Di-iso-decylphthalate(DIDP)		0.005	N.D.	N.D.	N.D.	---
Total (DINP+DNOP+DIDP)		---	< 0.015	< 0.015	< 0.015	< 0.1

Remark: *****The Limited value is based on EC Regulation 1907/2006, REACH Annex XVII (entry 51&52).

13. Bisphenol-A (BPA) Content

Test Method: With reference to EPA3550C-2007, Analysis was performed by UPLC-MS-MS.

Test Item	Unit	MDL	Test Results			Limited Value*****
			4#	5#	6#	
Bisphenol-A (BPA)	mg/kg	0.1	N.D	N.D	N.D	Prohibition

Remark: *****The Limited value is based on court case RG17859873, RG17880128.

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14. Phthalates Contents

Test Method: With reference to CPSC-CH-C1001-09.4(2018), Analysis was performed by GC-MS.

Test Item(s)		Results (%)			MDL (%)	Limited Value# (%)
		4#	5#	6#		
Di-(2-ethylhexyl) Phthalate	(DEHP)	N.D.	N.D.	N.D.	0.005	0.1
Dibutyl Phthalate	(DBP)	N.D.	N.D.	N.D.	0.005	0.1
Butylbenzyl Phthalate	(BBP)	N.D.	N.D.	N.D.	0.005	0.1
Di-iso-nonyl Phthalate	(DINP)	N.D.	N.D.	N.D.	0.010	0.1
Diisobutyl phthalate	(DIBP)	N.D.	N.D.	N.D.	0.005	0.1
Di-n-hexyl phthalate	(DHEXP/DnHP)	N.D.	N.D.	N.D.	0.005	0.1
Di-n-octyl Phthalate	(DNOP)	N.D.	N.D.	N.D.	0.005	0.1
Di-iso-decyl Phthalate	(DIDP)	N.D.	N.D.	N.D.	0.010	0.1

Remark: #The Limited value is based on US California Proposition 65& California Bill AB 1108:2007& Consumer Product Safety Improvement Act of 2008, CPSIA section 108.

15. Extractable Lead, Cadmium and Cobalt Content

Test Method: As specified in EEC Directive 84/500/EEC and 2005/31/EC, Analysis was ICP-MS.

Sample No. :1#-1	Result				Category 2 Limit
	1	2	3	4	
Released Lead (mg/l)	< 0.1	< 0.1	< 0.1	< 0.1	4.0
Released Cadmium (mg/l)	< 0.01	< 0.01	< 0.01	< 0.01	0.3
Released Cobalt (mg/l)	< 0.01	< 0.01	< 0.01	< 0.01	0.05
Volume of 4% Acetic acid used (ml)	380	380	380	380	---
Depth (mm)	86	86	86	86	---
Diameter (mm)	79	79	79	79	---

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Sample No. :2#-1	Result				Category 2 Limit
	1	2	3	4	
Released Lead (mg/l)	< 0.1	< 0.1	< 0.1	< 0.1	4.0
Released Cadmium (mg/l)	< 0.01	< 0.01	< 0.01	< 0.01	0.3
Released Cobalt (mg/l)	< 0.01	< 0.01	< 0.01	< 0.01	0.05
Volume of 4% Acetic acid used (ml)	380	380	380	380	---
Depth (mm)	85	85	85	85	---
Diameter (mm)	78	78	78	78	---

Sample No. :3#-1	Result				Category 2 Limit
	1	2	3	4	
Released Lead (mg/l)	< 0.1	< 0.1	< 0.1	< 0.1	4.0
Released Cadmium (mg/l)	< 0.01	< 0.01	< 0.01	< 0.01	0.3
Released Cobalt (mg/l)	< 0.01	< 0.01	< 0.01	< 0.01	0.05
Volume of 4% Acetic acid used (ml)	380	380	380	380	---
Depth (mm)	85	85	85	85	---
Diameter (mm)	78	78	78	78	---

Remark:

Category 1: Articles that cannot be filled and articles that can be filled, the internal depth of which does not exceed 25 mm.

Category 2: Articles, not in categories 1 or 3, which can be filled

Category 3: Cooking ware; packaging and storage vessels having a capacity of more than three liters

Lip and rim area for drinking utensils: With coating or colour within 20 mm along the lip and Rim area which is different from the colour of the interior cavity.

***** To be continued *****

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16. Sensorial Examination Odour and Taste Test

Test Method: Sensory test with reference to DIN 10955:2024-01

Test Medium: Distilled water

Test Condition: 70°C for 2 hours

Test Items	Result			Maximum Permissible Limit
	1#-1	2#-1	3#-1	
Sensorial examination odour (Point scale)	0	0	0	2.5
Sensorial examination taste (Point scale)	0	0	0	2.5

Scale evaluation:

- 0: No perceptible odour and taste
- 1: Odour and taste just perceptible (still difficult to define)
- 2: Moderate odour and taste
- 3: Moderately strong odour and taste
- 4: Strong odour and taste

17. Extractable Lead, Cadmium and Cobalt Content

Test Method: With reference to DIN EN 1388-1:1995 & DIN EN 1388-2:1995(Rim)-Determination of release of Lead and Cadmium from ceramic ware or silicate surfaces. Analysis was performed by ICP-MS

Sample No. :1#-1	Result				Hollow-ware Limit
	1	2	3	4	
Released Lead (mg/l)	< 0.1	< 0.1	< 0.1	< 0.1	See table 1
Released Cadmium (mg/l)	< 0.01	< 0.01	< 0.01	< 0.01	
Released Cobalt (mg/l)	< 0.01	< 0.01	< 0.01	< 0.01	0.1
Volume of 4% Acetic acid used (ml)	380	380	380	380	---
Depth (mm)	86	86	86	86	---
Diameter (mm)	79	79	79	79	---

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Sample No. :2#-1	Result				Hollow-ware Limit
	1	2	3	4	
Released Lead (mg/l)	< 0.1	< 0.1	< 0.1	< 0.1	See table 1
Released Cadmium (mg/l)	< 0.01	< 0.01	< 0.01	< 0.01	
Released Cobalt (mg/l)	< 0.01	< 0.01	< 0.01	< 0.01	0.1
Volume of 4% Acetic acid used (ml)	380	380	380	380	---
Depth (mm)	85	85	85	85	---
Diameter (mm)	78	78	78	78	---

Sample No. :3#-1	Result				Hollow-ware Limit
	1	2	3	4	
Released Lead (mg/l)	< 0.1	< 0.1	< 0.1	< 0.1	See table 1
Released Cadmium (mg/l)	< 0.01	< 0.01	< 0.01	< 0.01	
Released Cobalt (mg/l)	< 0.01	< 0.01	< 0.01	< 0.01	0.1
Volume of 4% Acetic acid used (ml)	380	380	380	380	---
Depth (mm)	85	85	85	85	---
Diameter (mm)	78	78	78	78	---

Sample No. :1#-2	Result				Rim Limit
	1	2	3	4	
Released Lead (mg/article)	< 0.1	< 0.1	< 0.1	< 0.1	2.0
Released Cadmium (mg/article)	< 0.01	< 0.01	< 0.01	< 0.01	0.2
Released Cobalt (mg/l)	< 0.01	< 0.01	< 0.01	< 0.01	0.1
Volume of 4% Acetic acid used (ml)	270	270	270	270	---
Depth (mm)	86	86	86	86	---
Diameter (mm)	79	79	79	79	---

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Sample No. :2#-2	Result				Rim Limit
	1	2	3	4	
Released Lead (mg/article)	< 0.1	< 0.1	< 0.1	< 0.1	2.0
Released Cadmium (mg/article)	< 0.01	< 0.01	< 0.01	< 0.01	0.2
Released Cobalt (mg/l)	< 0.01	< 0.01	< 0.01	< 0.01	0.1
Volume of 4% Acetic acid used (ml)	270	270	270	270	---
Depth (mm)	85	85	85	85	---
Diameter (mm)	78	78	78	78	---

Sample No. :3#-2	Result				Rim Limit
	1	2	3	4	
Released Lead (mg/article)	< 0.1	< 0.1	< 0.1	< 0.1	2.0
Released Cadmium (mg/article)	< 0.01	< 0.01	< 0.01	< 0.01	0.2
Released Cobalt (mg/l)	< 0.01	< 0.01	< 0.01	< 0.01	0.1
Volume of 4% Acetic acid used (ml)	270	270	270	270	---
Depth (mm)	85	85	85	85	---
Diameter (mm)	78	78	78	78	---

Permissible limit of Lead and Cadmium is quoted from DIN 51032:2017-07.

Table 1

Permissible limits for articles made from ceramics, glass ceramics with decorated inner surfaces.

Items	Flatware		Hollow-ware	
	Lead mg/dm ²	Cadmium mg/dm ²	Lead mg/l	Cadmium mg/l
Tableware Kitchen Equipment	0.8*)	0.07*)	4.0*)	0.3*)
Cooking & baking utensils, receptacles also used as packaging storage container	0.4	0.05	1.5*)	0.1*)

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*) In agreement with EC Directive

Table 2
Permissible limits for drinking rim

Lead, in mg per article	Cadmium, in mg per article
2.0	0.20

Test Part Description:

1#-1 Blue ceramic cup (inner wall)

2#-1 Green ceramic cup (inner wall)

3#-1 Grey ceramic cup (inner wall)

4# Blue coating

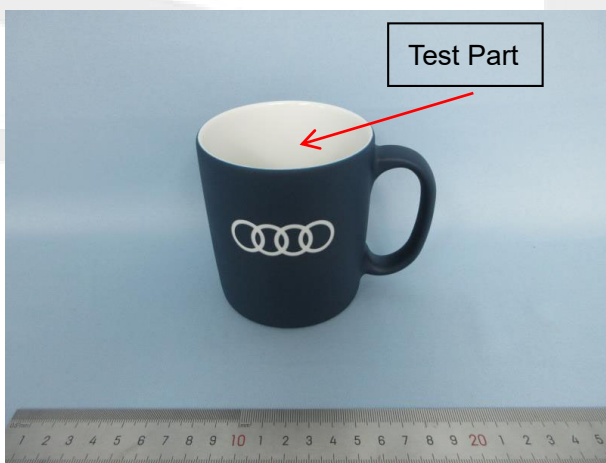
5# Green coating

6# Grey coating

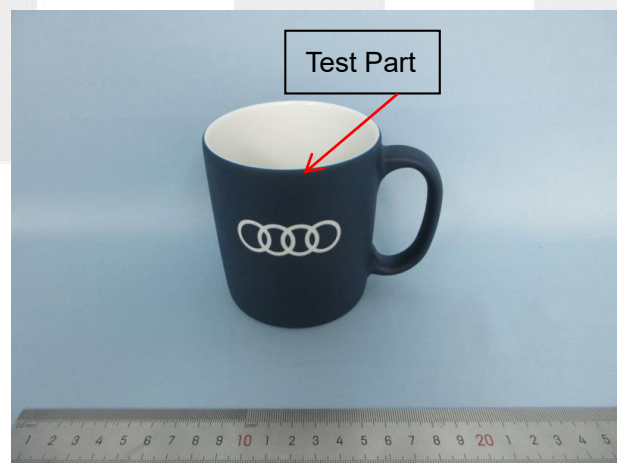
1#-2 Blue ceramic cup (inner wall) (rim)

2#-2 Green ceramic cup (rim)

3#-2 Grey ceramic cup (rim)

SAMPLE PHOTOS

1#-1



1#-2

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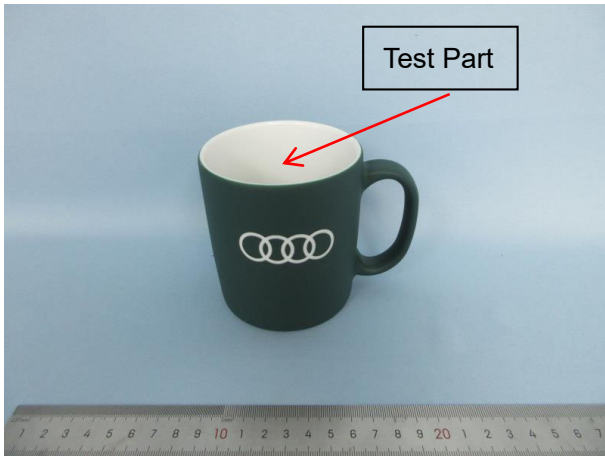
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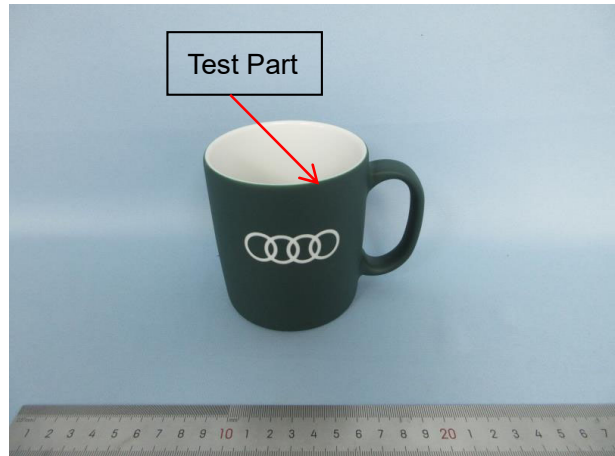
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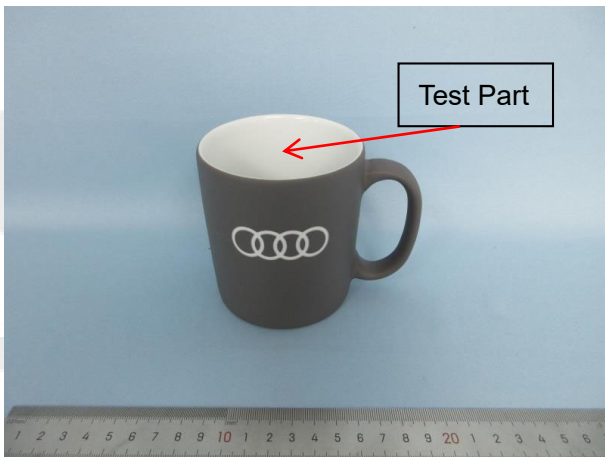
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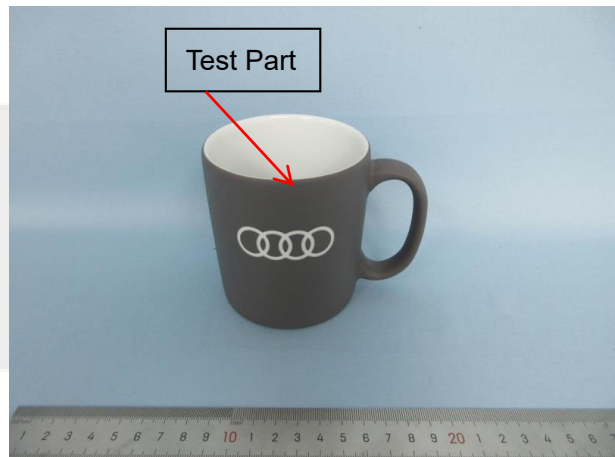
2#-1



2#-2



3#-1



3#-2



4#



5#

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GENERAL CONDITIONS OF SERVICES

STQ Testing Services Co.,Ltd. (hereinafter "STQ"), The testing or examining under the request of the customer should obey terms as follow, according to regulation of "Contract Law of the People's Republic of China" on processing and undertaking contract, our company have legal right of termination without any reason and have the right to accept or refuse testing or examining request:

1. STQ only acts for the person or body originating the instructions (the "Clients"). No other party is entitled to give instructions, particularly on the scope of testing or delivery of report or certificate, unless authorized by the Clients.
2. The delivery and return fee of the samples which need to do testing at STQ should be paid by the client. STQ will not bear the responsibility for the testing error that is caused by transporting, packaging and labelling.
3. Sample recycling: when the testing or examining is finished, the customer should recycle the sample. Within 30 days after issuing of testing report, if the customer could not recycle the sample or send notification of sample recycling in written (for example, if the sample belongs to consumables, toxic drugs, dangerous goods and other items that are not suitable for long-term storage, such as semi-finished products and fragile samples such as liquids and powders, the retention period will be shortened to 7 days). After the retention period, STQ has the right to dispose of the sample arbitrarily without paying compensation or compensation to the customer and take no responsibility for the consequences that damages the customer's trade secrets and intellectual property rights due to the loss of the sample.
4. The Clients shall always comply with the following before or during STQ providing its services:
 - a) provide sample(s) and relevant data, at the same time, guarantee the consistence of the sample(s)' name they declared with the sample(s) or the goods provided. Otherwise, STQ will not bear any relevant responsibilities;
 - b) giving timely instructions and adequate information to enable STQ to perform the services effectively;
 - c) supply, when requested by STQ, any equipment and personnel for the performance of the services;
 - d) take all necessary steps to eliminate or remedy any obstruction in the performance of the services;
 - e) inform STQ in advance of any hazards or dangers, actual or potential, associated with any order of samples or testing;
 - f) provide all necessary access for STQ's representative to enable the required services to be performed effectively;
 - g) ensure all essential steps are taken for safety of working conditions, sites and installations during the performance of services;
 - h) fully discharge all its liabilities under any contract like sales contract with a third party, whether or not a report or certificate has been issued by STQ, failing which STQ shall be under no obligation to the Clients.
5. Subject to STQ's accepting the Client's instructions, STQ will issue reports or certificates which reflect statements of opinion made with due care within the scope of instructions but STQ is not obliged to report upon any facts outside the instructions, if there were any dissidence about the report or certificate, the Client should provide the written declaration to STQ within 15 days after the date receiving the report or certificate, otherwise, STQ will not hear the case after the date limit.
6. STQ is irrevocably authorized by the Clients to deliver at its discretion the report or the certificate to any third party when instructed by the Clients or where it implicitly follows from circumstances, trade custom, usage or practice as determined by STQ.
7. A test report will be issued in confidence to the Clients and it will be strictly treated as such by STQ. 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 STQ. The Clients to whom the Report is issued may, however, show or send it, or a certified copy thereof prepared by STQ, to his customer, supplier or other persons directly concerned. STQ 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.
8. Applicants wishing to use STQ's reports in court proceedings or arbitration shall inform STQ to that effect prior to submitting the sample for testing.
9. The report will refer only to the sample tested and will not apply to the bulk, unless the sampling has been carried out by STQ and is stated as such in the Report. Also, the report is only for reference.
10. Any documents containing engagements between the Clients and third parties like contracts of sale, letters of credit, bills of lading, etc. are regarded as information for STQ only and do not affect the scope of the services or the obligations accepted by STQ.
11. If the Clients do not specify the methods/standards to be applied, STQ will choose the appropriate ones and further information regarding the methods can be obtained by direct contact with STQ, for the in-house method, STQ will only provide the summary.
12. No liability shall be incurred by and no claim shall be made against STQ or its servants, agents, employees or independent contractors in respect of any loss or damage to any such materials, equipment and property occurring whilst at STQ or any work places in which the testing is carried out, or in the course of transit to or from STQ or the said work places, whether or not resulting from any acts, neglect or default on the part of any such servants, agents, employees or independent contractors of STQ.
13. STQ will not be liable, or accept responsibility for any loss or damage howsoever arising from the use of information contained in any of its reports or in any communication whatsoever about its said tests or investigations.
14. Except for term 11 and term 12, if the test sample is damaged due to the negligence of STQ, the total compensation for loss and damage to the sample or loss to the customer shall not exceed twice of the test service fee.
15. In the event of STQ prevented by any cause outside STQ's control from performing any service for which an order has been given or an agreement made, the Clients shall pay to STQ:
 - a) the amount of all abortive expenditure actually made or incurred;

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- b) a proportion of the agreed fee or commission equal to the proportion (if any) of the service actually carried out by STQ, and STQ shall be relieved of all responsibility whatsoever for the partial or total non-performance of the required service.
16. STQ shall be discharged from all liabilities for all claims for loss, damage or expense unless suit is brought within one calendar year after the date of the performance by STQ of the service relating to the claim or in the event of any alleged non-performance within one year of the date when such service should have been completed.
17. The Clients acknowledge that STQ does not, either by entering into a contract or by performing service, assume or undertake to discharge any duty of the Clients to any other persons. STQ is neither an insurer nor a guarantor and disclaims all liability in such capacity.
18. The Clients shall hold harmless and indemnify STQ and its officers, employees, agents or independent contractors against all claims made by any third party for loss, damage or expense of whatsoever nature including reasonable legal expenses relating to the performance or non-performance of any services to the extent that the aggregate of any such claims relating to any one service exceed the limits mentioned in Clause 13.
19. Any unauthorized alteration, forgery or falsification of the content or appearance of the report/certificate is unlawful and offenders may be prosecuted to the fullest extent of the law; in the event of improper use of the report, STQ reserves the right to withdraw it, and to adopt any other measures which may be appropriate.
20. Samples are deposited with and accepted by STQ on the basis that either they are insured by the Clients or the Clients assumes entire responsibility for loss through fire, theft, burglary or for damages arising in the course of analysis or handling, without recourse whatsoever to STQ or its servants, agent, employees or independent contractors.
21. If the requirements of the Clients require the analysis of samples by the Clients' or any third party's laboratory, STQ will only convey the result of the analysis without responsibility for its accuracy. If STQ is only able to witness an analysis by the Clients' or any third Party's laboratory STQ will only confirm that the correct sample has been analyzed without responsibility for the accuracy of any analysis or results.
22. In the event of any unforeseen additional time or costs being incurred in the course of carrying out any of its services, STQ shall be entitled to charge the Clients additional fees to reflect the additional time and costs incurred.
23. All rights (including but not limited to copyright) in any reports, certificates or other materials produced by STQ in the course of providing its services shall remain vested in STQ.
24. Unless otherwise agreed in written, payment should be arranged within 10 days after the invoice date or the debit note date. If the payment is overdue, the overdue penalty shall be calculated at 1% per day of the unpaid part till the actual payment date. All expenses, costs and losses incurred by STQ as a result of collecting or claiming the fees owed shall be borne by the customer, including but not limited to attorney fees, litigation fees, preservation fees, preservation guarantee fees, travel expenses, etc.
25. Test results may be transmitted by electronic means at the Client's request. However, it should be noted that electronic transmission cannot guarantee the information contained will not be lost, delayed or intercepted by third party. STQ is not liable for any disclosure, error or omission in the content of such messages as a result of electronic transmission.
26. If necessary, STQ may subcontract part of or all tests to competent subcontractors. If no objection is raised at the time of the Clients submitting the application, STQ shall assume the Client's approval.
27. This report/certificate does not relieve sellers/suppliers from their contractual responsibility with regards to the quality/quantity of this delivery nor does it prejudice the Client's right to claim towards sellers/suppliers for compensation for any apparent and/or hidden defects not detected during STQ's random inspection or testing or audit.
28. The testing data and result(s) in this report is(are) just for scientific research, education, internal quality control and product development etc.
29. STQ reserves the right to include Special Conditions in addition to the foregoing General Conditions if warranted by the particular circumstances of the required test or investigation [this clause is only effective when the other party has been informed].
30. The foregoing General Conditions shall in all respects be governed, construed, interpreted and operated in accordance with the relevant Chinese laws and regulations. Unless otherwise agreed, the arbitration shall take place in P. R. C
31. These General Condition have been drafted in Chinese and may be translated into other languages. In the event of any discrepancy, the Chinese version shall prevail.
32. In general sample will be stored for 30 days. But for liquid, powder, etc semi-product & fragile product, it will be stored for 15 days after the report is issued.

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