

Test Report

Report No. : AGC03507210305-001

SAMPLE NAME : Foldable Chair with Cooler Bag

MODEL NAME : MO6112

APPLICANT: MID OCEAN BRANDS B.V

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

DATE OF ISSUE: Mar.15, 2021

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



Any report having not been signed by authorized approver, or having been altered without authorization, or having not been stamped by the coefficient estimated approver, or having been altered without authorization, or having not been stamped by the coefficient estimated approver, or having been altered without authorization, or having not been stamped by the coefficient estimated approver, or having been altered without authorization, or having not been stamped by the coefficient estimated in the report apply only to the tested sample. Any objections to report issued by AGC should be submitted to AGC within 15day after the issuance of the test report. Further enquiry of validity or verification of the test report should be addressed to AGC by agc@agc-cert.com.



Report No.: AGC03507210305-001

Page 1 of 7

Applicant : MID OCEAN BRANDS B.V

Address 7/F, Kings Tower, 111 King Lam Street, Cheung Sha Wan, Kowloon,

Hong Kong.

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 : Foldable Chair with Cooler Bag

Model : MO6112 Vendor Code : 107978 Sample Received Date : Mar.10, 2021

Testing Period : Mar.10, 2021 to Mar.15, 2021

Test Requested: Conclusion

1. As specified by client, to determine the Aromatic Amines Azodyes(AZO) content in

the submitted sample(s) with reference to entry 43, Annex XVII of the REACH Regulation (EC) No 1907/2006.

2. As specified by client, to determine the Phthalates content in the submitted sample(s) with reference to entry 51&52, Annex XVII of the REACH Regulation

(EC) No 1907/2006.
3. As specified by client, to determine the Cadmium(Cd) content in the submitted sample(s) with reference to entry 23, Annex XVII of the REACH Regulation (EC)

Pass

No 1907/2006.

4. As specified by client, to determine the Lead(Pb) content in the submitted sample(s) with reference to entry 63, Annex XVII of the REACH Regulation (EC) No

1907/2006.As specified by client, to determine the Color fastness to rubbing of the submitted sample.

Pass

Pass

Pass

Jossie-lians

Approved by: humgushua

Approved by:

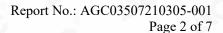
Huangguohua

Liangdan, Jessie.Liang

Vice Laboratory Manager

Technical Director

Any report having not been signed by authorized approver, or having been altered without authorization, or having not been stamped by the content of Stamp" is deemed to be invalid. Copying or excerpting portion of, or altering the content of the report is not permitted without the written authorization of AGC. The test results presented in the report apply only to the tested sample. Any objections to report issued by AGC should be submitted to AGC within 15days after the issuance of the test report. Further enquiry of validity or verification of the test report should be addressed to AGC by agc@agc-cert.com.



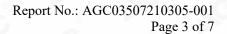


Test Result(s):

1. Test Result of Aromatic Amines Azodyes(AZO) Content

Test Item	Test Method/ Instrument	MDL	Limit
4-Aminobiphenyl(CAS No.:92-67-1)		5mg/kg	≤30mg/kg
Benzidine(CAS No.:92-87-5)	5 ⁰ 20	5mg/kg	≤30mg/kg
4-Chloro-o-Toluidine (CAS No.:95-69-2)		5mg/kg	≤30mg/kg
2-Naphthylamine(CAS No.: 91-59-8)		5mg/kg	≤30mg/kg
4-amino-2',3-dimethylazobenzene (CAS No.:97-56-3)	500 CC	5mg/kg	≤30mg/kg
5-Nitro-o-toluidine(CAS No.:99-55-8)	· ·	5mg/kg	≤30mg/kg
4-Chloroaniline(CAS No.:106-47-8)	. GO &	5mg/kg	≤30mg/kg
4-Methoxy-m-phenylenediamine (CAS No.: 615-05-4)		5mg/kg	≤30mg/kg
4,4'-Diaminodiphenylmethane (CAS No.: 101-77-9)		5mg/kg	≤30mg/kg
3,3'-Dichlorobenzidine (CAS No.: 91-94-1)		5mg/kg	≤30mg/kg
3,3'-Dimethoxybenzidine (CAS No.: 119-90-4)	EN ISO 14362-1:2017& EN ISO 14362-3:2017 a/	5mg/kg	≤30mg/kg
3,3'-Dimethybenzidine (CAS No.:119-93-7)	GC-MS	5mg/kg	≤30mg/kg
4,4'-Methylenedi-o-toluidine (CAS No.:838-88-0)		5mg/kg	≤30mg/kg
6-methoxy-m-toluidine (CAS No.:120-71-8)	NGC - C	5mg/kg	≤30mg/kg
4,4'-methylenebis[2-chloroaniline] (CAS No.:101-14-4)		5mg/kg	≤30mg/kg
4,4'-Oxydianiline(CAS No.:101-80-4)	-C	5mg/kg	≤30mg/kg
4,4'-Thiodianiline(CAS No.:139-65-1)	9 . 60	5mg/kg	≤30mg/kg
2-Aminotoluene(CAS No.:95-53-4)	。	5mg/kg	≤30mg/kg
4-methyl-m-phenylenediamine (CAS No.:95-80-7)		5mg/kg	≤30mg/kg
2,4,5-Trimethylaniline (CAS No.:137-17-7)	For Co	5mg/kg	≤30mg/kg
2-Methoxyaniline(CAS No.:90-04-0)	0	5mg/kg	≤30mg/kg
4-Aminoazobenzene (CAS No.:60-09-3)		5mg/kg	≤30mg/kg

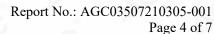
Any report having not been signed by authorized approver, or having been altered without authorization, or having not been stamped by the Bedicated Pesting/Inspection Stamp" is deemed to be invalid. Copying or excerpting portion of, or altering the content of the report is not permitted without the writter authorization of AGC. The test results presented in the report apply only to the tested sample. Any objections to report issued by AGC should be submitted to AGC within 15day after the issuance of the test report. Further enquiry of validity or verification of the test report should be addressed to AGC by agc@agc-cert.com.





T 414 (A)	6.0	Result(s)) (mg/kg)	
Test Item(s)	1-1	1-2	1-3	1-4
4-Aminobiphenyl	N.D.	N.D.	N.D.	N.D.
Benzidine	N.D.	N.D.	N.D.	N.D.
4-Chloro-o-Toluidine	N.D.	N.D.	N.D.	N.D.
2-Naphthylamine	N.D.	N.D.	N.D.	N.D.
4-amino-2',3-dimethylazobenzene	N.D.	N.D.	N.D.	N.D.
5-Nitro-o-toluidine	N.D.	N.D.	N.D.	N.D.
4-Chloroaniline	N.D.	N.D.	N.D.	N.D.
4-Methoxy-m-phenylenediamine	N.D.	N.D.	N.D.	N.D.
4,4'-Diaminodiphenylmethane	N.D.	N.D.	N.D.	N.D.
3,3'-Dichlorobenzidine	N.D.	N.D.	N.D.	N.D.
3,3'-Dimethoxybenzidine	N.D.	N.D.	N.D.	N.D.
3,3'-Dimethybenzidine	N.D.	N.D.	N.D.	N.D.
4,4'-Methylenedi-o-toluidine	N.D.	N.D.	N.D.	N.D.
6-methoxy-m-toluidine	N.D.	N.D.	N.D.	0 N.D.
4,4'-methylenebis[2-chloroaniline]	N.D.	N.D.	N.D.	N.D.
4,4'-Oxydianiline	N.D.	N.D.	N.D.	N.D.
4,4'-Thiodianiline	N.D.	N.D.	N.D.	N.D.
2-Aminotoluene	N.D.	N.D.	N.D.	N.D.
4-methyl-m-phenylenediamine	N.D.	N.D.	N.D.	N.D.
2,4,5-Trimethylaniline	N.D.	N.D.	N.D.	N.D.
2-Methoxyaniline	N.D.	N.D.	N.D.	N.D.
4-Aminoazobenzene	N.D.	N.D.	N.D.	N.D.
Conclusion	Conformity	Conformity	Conformity	Conformit

Any report having not been signed by authorized approver, or having been altered without authorization, or having not been stamped by the Bedicated Pesting/Inspection Stamp" is deemed to be invalid. Copying or excerpting portion of, or altering the content of the report is not permitted without the writter authorization of AGC. The test results presented in the report apply only to the tested sample. Any objections to report issued by AGC should be submitted to AGC within 15day after the issuance of the test report. Further enquiry of validity or verification of the test report should be addressed to AGC by agc@agc-cert.com.





2. Test Result of Phthalates Content

Test Item	Test Method/ Instrument	MDL	Limit
Diisobutyl phthalate(DIBP)		0.010%	8
(CAS No.: 84-69-5)		0.01070	©
Dibutyl phthalate (DBP)	8	0.010%	1 -0
(CAS No.: 84-74-2)		0.01076	Single<0.1%
Butylbenzyl phthalate (BBP)		0.010%	Sum<0.1%
(CAS No.: 85-68-7)		0.01076	@
Di-(2-ethylhexyl) Phthalate (DEHP)	TN 1 1250 200 1/ GG 3 50	0.010%	
(CAS No.: 117-81-7)	EN 14372:2004/ GC-MS	0.01070	
Di-n-octyl phthalate (DNOP)		0.010%	
(CAS No.: 117-84-0)	1 0 -0	0.01076	
Di-isononyl phthalate (DINP)		0.010%	G .0.10/
(CAS No.: 28553-12-0; 68515-48-0)		0.01070	Sum<0.1%
Di-isodecyl phthalate(DIDP)		0.0100/	
(CAS No.: 26761-40-0; 68515-49-1)		0.010%	

	Test result (%)						a.C			
No.	DIBP	DBP	BBP	DEHP	Sum(DIBP+DBP +BBP+DEHP)	DNOP	DINP	DIDP	Sum(DNOP+ DINP+DIDP)	Conclusion
1-5	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	Conformity
1-6	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	Conformity
1-7	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	Conformity

3. Test Result of Cadmium(Cd) Content

Test Item	Cadmium(Cd) (CAS No.: 7440-43-9)				
Limit(mg/kg)	<100				
MDL(mg/kg)	10				
Test Method/ Instrument	IEC 62321-5:2013/ ICP-OES				

	No.	Test result (mg/kg)	
ſ		Cadmium(Cd)	Conclusion
- G-	1-1	N.D.	Conformity
	1-2	N.D.	Conformity
	1-3	N.D.	Conformity
	1-4	N.D.	Conformity
	1-5	N.D.	Conformity
-C	1-6	N.D.	Conformity
	1-7	N.D.	Conformity

Any report having not been signed by authorized approver, or having been altered without authorization, or having not been stamped by the Bedicated Pesting/Inspection Stamp" is deemed to be invalid. Copying or excerpting portion of, or altering the content of the report is not permitted without the writter authorization of AGC. The test results presented in the report apply only to the tested sample. Any objections to report issued by AGC should be submitted to AGC within 15day after the issuance of the test report. Further enquiry of validity or verification of the test report should be addressed to AGC by agc@agc-cert.com.



Report No.: AGC03507210305-001 Page 5 of 7

NT.	Test result (mg/kg)	Gamalanian
No.	Cadmium(Cd)	Conclusion
1-8	N.D.	Conformity
1-9	N.D.	Conformity
1-10	N.D.	Conformity
1-11	N.D.	Conformity

4. Test Result of Lead(Pb) Content

Test Item	Lead(Pb) (CAS No.: 7439-92-1)				
Limit(mg/kg)	<500				
MDL(mg/kg)	10				
Test Method/ Instrument	IEC 62321-5:2013/ ICP-OES				

	.,	Test result (mg/kg)	
	No.	Lead(Pb)	Conclusion
	1-1	N.D.	Conformity
	1-2	N.D.	Conformity
	1-3	N.D.	Conformity
	1-4	N.D.	Conformity
- C	1-5	N.D.	Conformity
	1-6	N.D.	Conformity
8	1-7	N.D.	Conformity
-,0	1-8	N.D.	Conformity
	1-9	N.D.	Conformity
8	1-10	N.D.	Conformity
	1-11	N.D.	Conformity

Note:

mg/kg = milligram per kilogram N.D.=Not Detected (less than method detection limit)

MDL = Method Detection Limit %= percentage

Any report having not been signed by authorized approver, or having been altered without authorization, or having not been stamped by the Bedicated Pesting/Inspection Stamp" is deemed to be invalid. Copying or excerpting portion of, or altering the content of the report is not permitted without the written appropriation of AGE. The test results presented in the report apply only to the tested sample. Any objections to report issued by AGC should be submitted to AGC within 15day after the issuance of the test report. Further enquiry of validity or verification of the test report should be addressed to AGC by agc@agc-cert.com.



Remark:

- As specified by client, only test the designated sample.
- ^a The EN ISO 14362-1:2017 methods will enable further cleavage of 4-aminoazobenzene to non-forbidden amines: aniline and 1,4-phenylenediamine, therefore, the test method of EN ISO 14362-3:2017 was employed to verify the presence of 4-aminoazobenzene

5. Test Result of Color fastness to rubbing

Test Item	Test Method		ult(s) -1	Limit (Client's Requirement)
		dry	wet	- Kequirement)
Color fastness to rubbing (Grade)	ISO 105-X12:2016	4-5	4-5	2-3
Conclusion		Conformity	Conformity	/

Test Item	Test Method		ult(s) -2	Limit (Client's Requirement)
0	100	dry	wet	Requirement
Color fastness to rubbing (Grade)	ISO 105-X12:2016	4-5	4-5	2-3
Conclusion	9 10	Conformity	Conformity	

Test Item	Test Method		Result(s)		
Test Item	rest Method	dry wet		Requirement)	
Color fastness to rubbing (Grade)	ISO 105-X12:2016	4-5	4-5	2-3	
Conclusion	1 20	Conformity	Conformity	120	

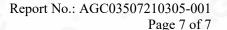
Note:

- Color fastness grade: grey scale (5 grade is good, 1 grade is bad).

Sample Description:

1-1	Blue cloth
1-2	Black cloth
1-3	Black hand strap
1-4	Black zipper cloth
1-5	Black zipper teeth
1-6	Black plastic strip
1-7	White plastic

Any report having not been signed by authorized approver, or having been altered without authorization, or having not been stamped by the Dedicated Festiva/Inspection Stamp" is deemed to be invalid. Copying or excerpting portion of, or altering the content of the report is not permitted without the written authorization of AGC where the test results presented in the report apply only to the tested sample. Any objections to report issued by AGC should be submitted to AGC within 15day after the issuance of the test report. Further enquiry of validity or verification of the test report should be addressed to AGC by agc@agc~cert.com.





1-8	Aluminum foil)		10	\G	
1-9	Metal zipper pull	< GO	-6	8		10
1-10	Metal rack		10	10°		@
1-11	Metal nail		0		10	-60

Test Flow Chart

1. For AZO, Phthalates Cutting/Preparation Weigh Sample Sample solvent extraction Concentration/ Dilution of Extracted solution GC-MS Filtration **DATA** 2. For Cd, Pb Acid digestion with Sample Preparation Weigh Sample microwave/hotplate Filtration DATA **ICP-OES**

The photo of the sample



AGC authenticate the photo only on original report

*** End of Report ***

Any report having not been signed by authorized approver, or having been altered without authorization, or having not been stamped by the Bedicated Restriction Stamp" is deemed to be invalid. Copying or excerpting portion of, or altering the content of the report is not permitted without the writter exphorization of AGC. The test results presented in the report apply only to the tested sample. Any objections to report issued by AGC should be submitted to AGC within 15day after the issuance of the test report. Further enquiry of validity or verification of the test report should be addressed to AGC by agc@agc-cert.com.



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.

Any report having not been signed by authorized approver, or having been altered without authorization, or having not been stamped by the Bedicated Pestud/Inspection Stamp" is deemed to be invalid. Copying or excerpting portion of, or altering the content of the report is not permitted without the written authorization of AGC. The test results presented in the report apply only to the tested sample. Any objections to report issued by AGC should be submitted to AGC within 15days after the issuance of the test report. Further enquiry of validity or verification of the test report should be addressed to AGC by agc@agc-cert.com.