

# **Test Report**

Report No. : AGC05443230406-001

- SAMPLE NAME : Anti-bacterial pen
- MODEL NAME : MO9951
- APPLICANT : MID OCEAN BRANDS B.V
- **STANDARD(S)** : Please refer to the following page(s).
- **DATE OF ISSUE** : May 12, 2023







#### MID OCEAN BRANDS B.V

Report No.: AGC05443230406-001

: 7/F, Kings Tower, 111 King Lam Street, Cheung Sha Wan, Kowloon, Hong Kong.

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

:

:

Sample Name	:	Anti-bacterial pen
Model	:	MO9951
Vendor code	:	108694
Country of Origin	:	CHINA
Country of Destination	:	EUROPE
Sample Received Date	:	Apr. 11, 2023
Testing Period	:	Apr. 11, 2023 to May 11, 2023
Test Requested	:	Selected test(s) as requested by client.

Test Requested:	Conclusion
Annex XVII of the REACH Regulation (EC) No 1907/2006, entry 63 - Lead(Pb) Content	Pass
Annex XVII of the REACH Regulation (EC) No 1907/2006, entry 23 -Cadmium(Cd) Content	Pass
Annex XVII of the REACH Regulation (EC) No 1907/2006, entry 27 - Nickel Release	Pass
-Antibacterial activity of Escherichia coli, Staphylococcus aureus and Candida albicans	See the result(s)

Approved by : Jessie ling

Liangdan, Jessie.Liang

**Technical Director** 



#### Report No.: AGC05443230406-001

Report Revise Record							
Report Version	Issued Date	Valid Version	Notes				
/	May 12, 2023	Valid	Initial release				



# The photo of the sample



The photo of AGC05443230406-001 is for use only with the original report.

Test I onte Description	
Test point	Test point description
1-1	White plastic pen holder
1-2	Plastic pen holder tip+Plastic pen holder head+Plastic pen middle ring
1-3	White plastic refill pen tube
1-4	Metal pen clip
1-5	Metal spring
1-6	Blue ink

# **Test Point Description**



Note: N.D.=Not Detected (less than method detection limit), MDL = Method Detection Limit, 1mg/kg=0.0001%

#### Annex XVII of the REACH Regulation (EC) No 1907/2006, entry 63

#### - Lead(Pb) Content

Test Methods and Equipment: IEC 62321-5:2013; ICP-OES

Test Item(s)	Unit Limit		MDL	Test Result(s)		
Test Item(s)		LIIIII	MDL	1-1	1-2	1-3
Lead(Pb)	mg/kg	500	10	N.D.	N.D.	N.D.
Con	Conformity	Conformity	Conformity			

Test Item(s)	Unit Limit		MDL	Test Result(s)		
Test Item(s)		LIIIII	WIDL	1-4	1-5	1-6
Lead(Pb)	mg/kg	500	10	N.D.	N.D.	N.D.
Conclusion				Conformity	Conformity	Conformity

Remark:

1. As specified by client, the submitted samples were mixed to test, the test points: 1-2

2. As specified by client, the submitted samples were directly tested without drying, the test points: 1-6

#### Annex XVII of the REACH Regulation (EC) No 1907/2006, entry 23

#### -Cadmium(Cd) Content

Test Methods and Equipment: IEC 62321-5:2013; ICP-OES

Test Item(s)	Unit	Limit	MDI	Test Result(s)		
Test Item(s)	Unit	Limit	MDL	1-1	1-2	
Cadmium(Cd)	mg/kg	100	10	N.D.	N.D.	
Conclusion				Conformity	Conformity	

Test Item(s)	Unit	Limit	MDL	Test Result(s)		
Test Item(s)	Unit	Liiiit	MDL	1-3	1-6	
Cadmium(Cd)	mg/kg	100	10	N.D.	N.D.	
Conclusion				Conformity	Conformity	

Remark:

1. As specified by client, the submitted samples were mixed to test, the test points: 1-2

2. As specified by client, the submitted samples were directly tested without drying, the test points: 1-6



### Annex XVII of the REACH Regulation (EC) No 1907/2006, entry 27

- Nickel Release

Test Methods and Equipment: EN 12472:2020 & EN 1811:2011+A1:2015; ICP-OES

Test Point(s)	Parallel Sample	Unit	Limit	MDL	Test Result(s) Nickel Release	Conclusion
	А	µg/cm²/week	0.5	0.05	N.D.	
1-4	В	µg/cm²/week	0.5	0.05	N.D.	Conformity
	С	µg/cm²/week	0.5	0.05	N.D.	

#### Limit requirements of Nickel Release

Nickel Release					
Type of sample	Pass	Fail			
Article with Nickel release limit of 0.5µg/cm <sup>2</sup> /week (Non-body piercing)	<0.88µg/cm <sup>2</sup> /week	≥0.88µg/cm <sup>2</sup> /week			
Article with Nickel release limit of 0.2µg/cm <sup>2</sup> /week (Body piercing)	<0.35µg/cm <sup>2</sup> /week	≥0.35µg/cm <sup>2</sup> /week			

#### - Antibacterial activity of Escherichia coli, Staphylococcus aureus and Candida albicans

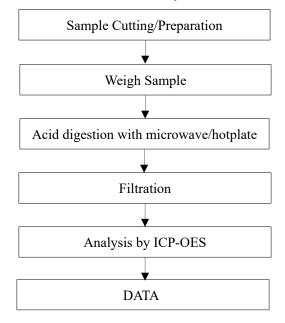
Test Point(s): 1-1

Test Method: ISO 22196:2011.

Test instrument: Biochemical incubator.

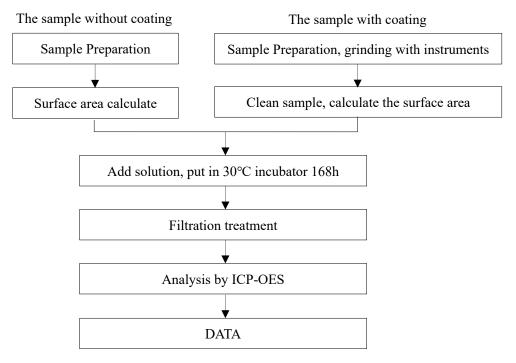
	Test strains					
	Escherichia coli ATCC 8739	Staphylococcus aureus ATCC 6538P	Candida albicans ATCC 10231			
Liquid inoculant concentration (each/mL)	5.1×10 <sup>5</sup>	4.2×10 <sup>5</sup>	3.3×10 <sup>5</sup>			
Liquid inoculant amount (mL)	0.4	0.4	0.4			
Untreated sample piece 0 h after inoculation on the number of living bacterium	1.5×10 <sup>4</sup>	1.5×10 <sup>4</sup>	1.3×10 <sup>4</sup>			
Untreated sample piece cultivate 24 h after inoculation on the number of living bacterium	$2.7 \times 10^{6}$	2.6×10 <sup>6</sup>	2.0×10 <sup>6</sup>			
Antibacterial sample piece cultivate 24 h after inoculation on the number of living bacterium	5.9×10 <sup>4</sup>	1.8×10 <sup>5</sup>	1.6×10 <sup>5</sup>			
Antibacterial activity value	1.66	1.16	1.10			
Antibacterial rate (%)	97.81	93.08	92.00			





## **Test Flow Chart of Heavy Metal Content**

## **Test Flow Chart of Nickel Release**





# 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. 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.

5. 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.

6. 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. 7. 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.

8. 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.
9. 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.

\*\*\* End of Report \*\*\*