

## Test Report

Number: GZHH00366861

Applicant: JoCo Sales & Marketing Inc.  
12100 Blue Valley Parkway, Overland Park,  
Kansas 66213

Date: Jun 18, 2020

### Sample Description:

One (1) style of submitted sample said to be :  
Item Name : **GSD ANTIBACTERIAL 80 WET WIPES.**  
Item Quantity : 5.  
Date Sample Received : Jun 04, 2020



### Ingredients List:

No.	CAS #	INGREDIENTS	WEIGHT %
1	8001-54-5	Benzalkonium Chloride	0.10%
2	123-03-5	HexadecylPyridinium Chloride	0.05%
3	104-29-0	Chlorphenesin	0.05%
4	57-55-6	Propanediol	0.20%
5	1406-18-4	Vitamin E	0.05%
6	85507-69-3	Aloe Vera Extract	0.10%
7	107-43-7	Amino acid moisturizer	0.10%
8	77-92-9	Citric acid	0.02%
9	56-81-5	Glycerol	0.50%
10	1968/4/2	Sodium citrate	0.02%
11	8001-54-5	Water	98.81%

\*\*\*\*\*

### Tests conducted:

As requested by the applicant, refer to attached page(s) for details.

\*\*\*\*\*

Intertek GM Testing Service Zhuhai Co. Ltd.

Sarah Xu  
Asst. Manager  
Healthcare and Beauty Products



## Test Report

Number: GZHH00366861

### Tests Conducted

1 Quantitative suspension test for evaluation of bactericidal activity of chemical disinfectants and antiseptics

With reference to BS EN 1276:2019

Dilution recommended for use:	No dilution
Product test concentration:	80% v/v
Active ingredient in product:	Benzalkonium chloride
Appearance:	Transparent colorless liquid
Contact time:	60s±5s
Test temperature:	(20±1) °C
Interfering substance:	0.3g/L bovine albumin (clean condition)
Inhibition method:	Dilution-neutralization
Neutralizing solution:	D/E neutralizing broth
Incubation:	(37±1) °C, 48 hours
Agar medium:	Trypticase Soy Agar
Test culture:	Escherichia coli K12 (NCTC 10538) Pseudomonas aeruginosa (ATCC 15442) Staphylococcus aureus (ATCC 6538) Enterococcus hirae (ATCC 10541) Severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2)

### Controls & validation:

Test microorganism	Validation suspension (cfu/ml) $N_v$ $N_{v0}=1/10N_v$ Criteria: $300 \leq N_v \leq 1600$	Experimental conditions control (cfu/ml) A Criteria: $A \geq 0.5 N_{v0}$	Neutralizer control (cfu/ml) B Criteria: $B \geq 0.5 N_{v0}$	Method validation (cfu/ml) C Criteria: $C \geq 0.5 N_{v0}$	Validity
Escherichia coli K12 (NCTC 10538)	950	91	88	97	Valid
Pseudomonas aeruginosa (ATCC 15442)	800	75	78	86	Valid
Staphylococcus aureus (ATCC 6538)	1000	88	89	96	Valid
Enterococcus hirae (ATCC 10541)	1000	78	85	97	Valid
Severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2)	900	95	96	90	Valid

\*\*\*\*\*



## Test Report

Number: GZHH00366861

### Tests Conducted

Result:

Test microorganism	Initial suspension(N) (cfu/ml) N <sub>0</sub> =1/10N Criteria: 1.5x10 <sup>8</sup> ≤N≤5.0x10 <sup>8</sup>	Final count (cfu/ml) N <sub>a</sub>	R(Log <sub>10</sub> Reduction) =Log N <sub>0</sub> -Log N <sub>a</sub>	%Reduction
Escherichia coli K12 (NCTC 10538)	4.1x10 <sup>8</sup>	<140	>5.0	>99.999
Pseudomonas aeruginosa (ATCC 15442)	1.8 x10 <sup>8</sup>	4.3 x10 <sup>3</sup>	3.6	99.976
Staphylococcus aureus (ATCC 6538)	4.2 x10 <sup>8</sup>	<140	>5.0	>99.999
Enterococcus hirae (ATCC 10541)	2.1 x10 <sup>8</sup>	<140	>5.0	>99.999
Severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2)	4.8 x10 <sup>8</sup>	4.1 x10 <sup>3</sup>	>5.0	>99.986

Remark:

- N = Test suspension, Number of cells per ml in bacterial suspensions.
- N<sub>0</sub> = (N<sub>0</sub>=1/10N), Number of cells per ml in the test mixtures at the beginning of the contact time (time 0).
- N<sub>a</sub> = Number of survivors per ml in the test mixtures at the end of the contact time.
- N<sub>v</sub> = Validation suspension, Number of cells per ml in bacterial suspensions.
- N<sub>v0</sub> = (N<sub>v0</sub> =1/10N<sub>v</sub>), Number of cells per ml in the test mixtures at the beginning of the contact time (time 0).
- A,B,C = Represent the different control test mixtures, A(experimental conditions control), B(Neutralizer control), C(Method validation)

Sample received condition: sample in unopened original

package. Date sample received: Jun 04, 2020

Testing period: Jun 04, 2020 to Jun 17, 2020

\*\*\*\*\*

End of report

*This report is made solely on the basis of your instructions and/or information and materials supplied by you. It is not intended to be a recommendation for any particular course of action. Intertek does not accept a duty of care or any other responsibility to any person other than the Client in respect of this report and only accepts liability to the Client insofar as is expressly contained in the terms and conditions governing Intertek's provision of services to you. Intertek makes no warranties or representations either express or implied with respect to this report save as provided for in those terms and conditions. We have aimed to conduct the Review on a diligent and careful basis and we do not accept any liability to you for any loss arising out of or in connection with this report, in contract, tort, by statute or otherwise, except in the event of our gross negligence or wilful misconduct. This report shall not be reproduced unless with prior written approval from Intertek GM Testing Services Zhuhai Co.,Ltd.*

