



TEST REPORT

Report No.: GXX24120158B(E)

Applicant: CAN-AMERICA PHMGH INC.2021

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Dllte :2025/01/24

Name of Sillnple	Polyhexamethylene guanidine effervescent tablet	Brand	GERMFAGA
Sample No.	GXX24120158	Sample appearance	Solid
Model	I	Sample Quantity	14 pieces
Date	December 2023	Blitth No.	202312501
Quality guarantee period	I	Test Type	Commission
Sample Received	2024/12/26	Test Period	2025/01/10-2025/01/22
Manufacturer	Jia'an Yujie (Beijing) Tracie Development Co., Ltd		
Address	No. 7167, Unit I, Building I, No.33 Guangshun North Street, Chaoyang District, Beijing		
Test Item	Coronavirus suspension quantitative inactivation test (including neutralizer identification test)		
Test Method	Technical Standard For disinfection(2002)		
Test Result/Conclusion	Please refer to next page(s)		
Note	I		

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I.The results of neutralizer identification test:

Neutralizer: 3.9% D/E neutralizing broth medium. Test strain: Human Coronavirus 229E,VR-740. Test cell lines: Human hepatocellular carcinoma cell line I-luh-7. Conditions of action: Acid I samples to 1000 mL of water, dissolve and shake well as the test solution, 5 min. Test method: Technical Standard For disinfection(2002) 2.1.1.10.			
Group Number	The logarithm of virus titer (TCID ₅₀ / mL) in each group of three experiments.		
	1	2	3
1	3.92	3.58	3.71
2	4.32	4.05	4.32
3	5.32	5.17	5.35
4	5.32	5.16	5.29
5	5.44	5.24	5.42
6	0	0	0

2. Coronavirus suspension quantitative inactivation test

Neutralizer: 3.9% D/E neutralizing broth medium. Test strain: Human Coronavirus 229E,YR-740. Test cell lines: Human hepatocellular carcinoma cell line Huh-7. Conditions of action: Add I samples to 1000 mL of water, dissolve and shake well as the test solution, 5 min. Test method: Technical Standard For disinfection(2002) 2.1.1.10.					
Test time	Group	The logarithm of virus titer (TCID ₅₀ / mL)		Killing log value	Killing rate (%)
		Control	Sample		
5 min	1	5.71	4.52	1.19	93.45
	2	5.42	4.29	1.13	92.68
	3	5.21	4.20	1.01	90.25

Note:The negative control cells grew well without cytopathic effect..

***** **END OF REPORT*******

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4. This report is invalid if being supplemented, deleted or altered.
5. Without written permission of our Company, this report can not be reproduced in part (except in whole).
6. The result(s) shown in this report refer only to the sample(s) tested.
7. Objection to this report must be submitted to our Company within 15 day . Otherwise, it will automatically deem to have accepted this report.
8. The Client shall be responsible for the accuracy, authenticity and completeness of the samples and information submitted for inspection, and the disputes arising therefrom shall be borne by the Client.
9. As any reports is issued as a result of this application for testing services, our Company will strictly keep confidentiality to the Clients. Except where disclosure is required on the basis of laws, regulations, judgments, and rulings (including in accordance with summons, court, or government proceedings).
10. The result(s) or conclusion(s) shown in this report about the description of the characteristics, composition, properties or quality are based on the specific time, methods and applicable criteria. Using different methods and criteria or under different environmental conditions for testing may come to different conclusions.
11. Since our Company's causes lead to modify the contents of this report, our Company shall reissue this report and bear the modification cost. The Client shall return the original report. Since the Client's causes lead to modify the contents of this report, the Client need to submit an application form for the reissue of report to our Company. The Client shall bear the modification cost and return the original report, if our Company approves to reissue this report.
12. The English version of this statement is translated from the Chinese one. If there is any disagreement between them, the Chinese version will be the final explanation.