

SUPER LABORATORY CO., LTD. TESTING CENTER TEST REPORT



CUSTOMER

Ji-Feng Biotechnology Co., Ltd.

7F, No. 446, Sec. 2, Zhongshan Rd., Zhonghe Dist., New Taipei City 235029, Taiwan (R.O.C.)

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Sample name : T-Pro Aqua EZ Clean

Manufacturer : Ji-Feng B	liotechnology C	o., Ltd.					
Manufacture date : 2024	/07		Sample expiry date : 2	2026/07			
Lot number : 900010107	1307		Country of origin : Tai	wan			
Sample condition : Chilled storage			Test sample packagin	Test sample packaging : Intact packaging			
Sample delivering : By a	pplicant		Number of samples :	58 g			
Contact person: Benny Sun			Report purpose : For	Report purpose : For export, Self-Management			
Phone number : (02)828	3-6236						
The sample inform	nation provided	above was cor	nfirmed by the customer				
Sample Received : 2024/09/20 Date of Test		esting: 2024/09/23	Date of Report : 2024/10/07				
Item	Result	Unit	Method	LOQ / LOD			
Antimicrobial activity : Pseudomonas aeruginos	97.95 a	%	Refer to ASTM E2315, Assessment of Antimicr				

----- Null below ------

Remarks :

• All inspection contents in this inspection report are inspected according to the entrusted items. If there is any falsehood, we are willing to take full responsibility.

a Time-Kill Procedure.

- Antimicrobial Activity: Please see the test appendix for detailed inspection results.
- Please note if the tests do not involve sampling, then the test report is only responsible for the test sample
 provided by the customer. And the report is invalid if not presented in full and for reference only; it shall not be
 used for advertising, sales promotions, or notarial purposes. Also, when the target is below the limit of detection
 (LOD) or the limit of quantification (LOQ), the test results will be expressed as "Negative" or "ND" (Not detected).



This report is used only for providing the testing results of commissioned items, not for determining the legality of the product.

Junh- ting frai SL Yueh-ting Tsai, Ph. D.

Approval Signatory



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Sample name : T-Pro Aqua EZ Clean Sample condition : Chilled storage Sample Received : 2024/09/20

Date of Testing : 2024/09/23

Date of Report : 2024/10/07



M61-240901415EN





Junh-ting frai Yueh-ting Tsai, Ph. D. Approval Signatory



Appendix

1. Specimens I.D. : M61-240901415

2. Test method :

2.1 Test strains and culture condition :

Test strains	Test condition	Culture condition	
Pseudomonas aeruginosa	25 ± 2°C	35 ± 2°C	
ATCC 9027	24 hr	48 ± 2 hr	

2.2 Reference :

2.2.1 Refer to ASTM E2315-23, Standard Guide for Assessment of Antimicrobial Activity Using a Time-Kill Procedure.

2.3 Grouping :

2.3.1 Test group : Test substances provided by customers.(After diluting 200 times with sterile water as specified by the customer, then perform the test.) 2.3.2 Control group : 0.85 % Sterile saline.

2.4 Prepare the suspension of test microbe, and the concentration is around $1.0 \times 10^8 \sim 1.0 \times 10^9$ CFU/mL.

2.5 Add 0.1 mL of microbial suspension into 10 mL test and control group.

2.6 The test groups were executed according to the conditions of 2.1.

2.7 Test population control groups were conducted for 10-fold serial dilution with 9 mL SCDLP broth at time zero, representing the concentration of the test organism present.

2.8 After serial dilution, each group was inoculated on appropriate medium. The medium was cultured under the culture condition. The growth was observed and colony counts were recorded.

2.9 Calculation :

LR = Log(A) - Log(D)

Reduction rate (R) = 100 % × (1 - 10^{-LR})

A = The number of microbe recovered from the test population control groups at time zero after inoculation.

D = The number of microbe recovered from the test groups incubated after the desired contact period.



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Appendix

3. Result :



5 Figure 2: Result of control and test group.

Figure 1: Test article.

Strains : Pseudomonas aeruginosa

P. aeruginosa M61-240901415 10³

Table 1 : Result of test article against test strain.

Test strains	Final inoculated concentration	Unit -	The residual amount of microbe			Reduction
Test strains	(CFU/mL)		Test population control groups (A)	Test groups (D)	LR	rate(%)1
	(CI O/IIIE)		(A)			
Pseudomonas aeruginosa	sa 7.1x10 ⁶	CFU/mL	8.4×10 ⁶	1.7×10 ⁵	1.69	97.95
i seudomonas deluginosa		Log	6.92	5.23		

1. Reduction rate (R) : If antimicrobial rate < 0.00 % presented with 0.00 %.

Table 2 : Conditions for the establishment of the test.

Condition ¹	Calculation result	Result	
The difference between the number of microbe washed immediately (A) in the control group and the number of microbe washed after the control group (B) was within the range of \pm 0.5 Log	Pseudomonas aeruginosa	0.44	-

1. Conditions for the establishment : All of them should be Pass. (Except when the test condition is more than 1 hour).