

[Cat. No.]      **K-6839**

### Introduction

*AccuPower*<sup>®</sup> *Pseudomonas aeruginosa* Real-Time PCR Kit is a product that can specifically detect *Pseudomonas aeruginosa* (*P. aeruginosa*) by real-time PCR.

*P. aeruginosa* is a gram-negative bacterium known to primarily cause sepsis, pneumonia and endocarditis. *P. aeruginosa* is the most common pathogen found in hospitalized patients of respiratory infections. There are more cases of acute infections in communities or hospitals than chronic infections, and in particular, infections in hospitals are often reported with antibiotic resistance.

This product contains all Real-time PCR components specific to *P. aeruginosa*, including DNA polymerase, dNTPs, and reaction buffer. The users can easily prepare reaction mixture simply by adding template DNA, Oligo Mix and DEPC-D.W.

### Features & Benefits

- Convenience: All necessary reactants for real-time PCR are included in a tube (i.e., Master Mix Type), allowing the users to perform reaction simply by adding template DNA, Oligo Mix, and DEPC-D.W.
- Sensitivity: By using BIONEER's HotStart *Taq* DNA Polymerase that minimizes non-specific reactions and maximizes reaction efficiency, only the target gene can be effectively amplified even with a trace amount of template DNA.

### Components

Components	Amount
2X Master Mix	625 µl x 2 ea
Oligo Mix	500 µl
DEPC-D.W.	1.8 ml
Positive Control (1x10 <sup>8</sup> copies/µl)	50 µl

\* **Note:** For research use only. Not for use in diagnostic or therapeutic procedures.

### Composition

Composition	25 µl reaction	
2X Master Mix	<i>Taq</i> DNA Polymerase	2.5 U
	dNTPs (dATP, dCTP, dGTP, dTTP)	Each 300 µM
	Reaction buffer with 2 mM MgCl <sub>2</sub>	1X
Oligo Mix	<i>P. aeruginosa</i> Forward primer	0.8 µM
	<i>P. aeruginosa</i> Reverse primer	0.8 µM
	<i>P. aeruginosa</i> Probe (FAM)	0.8 µM
	ROX dye	1X

### Specifications

<i>Taq</i> DNA Polymerase	
5'→3' exonuclease activity	Yes
3'→5' exonuclease activity	No
3'-A overhang	Yes

### Storage

Store at -20°C. If stored in the recommended temperature, this product will be stable until the expiration date printed out on the label.

### Online Resources



English

Visit our **product page** for additional information and protocols

### Ordering Information

Description	Cat. No.
<i>AccuPower</i> <sup>®</sup> <i>Pseudomonas aeruginosa</i> Real-Time PCR Kit, 1.25 ml of 2X Master Mix solution, 100 tests	K-6839




### Notice

BIONEER corporation reserves the right to make corrections, modifications, improvements and other changes to its products, services, specifications or product descriptions at any time without notice.

### Explanation of Symbols

<b>LOT</b> Batch Code	Biological Risks	<b>REF</b> Catalog Number	Caution
Consult Instructions For Use	Contains Sufficient for <n> tests	Do not Re-use	Manufacturer
<b>RUO</b> Research Use Only	Temperature Limitation	Use-by Date	

### Experimental Procedures

Steps		Procedure Details															
1	 <b>Preparation of reaction mixture</b>	1. Thaw all components of <i>AccuPower</i> <sup>®</sup> <i>Pseudomonas aeruginosa</i> Real-Time PCR Kit on ice and mix thoroughly before use. Then, briefly spin down all components.															
2	 <b>Composition of reaction mixture</b>	2. Add all components into PCR tubes (not provided) or a plate (not provided) referring to the following list of components (Based on 1 test). <table border="1" style="width: 100%; margin-top: 10px;"> <thead> <tr> <th style="text-align: left;">Components</th> <th style="text-align: left;">Amount</th> </tr> </thead> <tbody> <tr> <td>2X Master Mix</td> <td>12.5 µl</td> </tr> <tr> <td>Oligo Mix</td> <td>5 µl</td> </tr> <tr> <td>Template DNA</td> <td>1-5 µl</td> </tr> <tr> <td>DEPC-D.W.</td> <td>Variable</td> </tr> <tr> <td>Total volume</td> <td>25 µl</td> </tr> </tbody> </table>	Components	Amount	2X Master Mix	12.5 µl	Oligo Mix	5 µl	Template DNA	1-5 µl	DEPC-D.W.	Variable	Total volume	25 µl			
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3	 <b>Real-time PCR</b>	3. Place PCR tubes or plate on the Real-Time Quantitative thermal cycler. 4. Perform the reaction under the following conditions. <table border="1" style="width: 100%; margin-top: 10px;"> <thead> <tr> <th style="text-align: left;">Step</th> <th style="text-align: left;">Temperature</th> <th style="text-align: left;">Time</th> <th style="text-align: left;">Cycles</th> </tr> </thead> <tbody> <tr> <td>Pre-denaturation</td> <td>95°C</td> <td>5 min</td> <td>1 cycle</td> </tr> <tr> <td>Denaturation</td> <td>95°C</td> <td>10 sec</td> <td rowspan="2">45 cycles</td> </tr> <tr> <td>Annealing &amp; Extension</td> <td>55°C</td> <td>20 sec</td> </tr> </tbody> </table> <p>* <b>Note:</b> Users can adjust the protocol according to their instrument and template sequences to get optimal results.</p> 5. After the reaction is completed, analyze the results.	Step	Temperature	Time	Cycles	Pre-denaturation	95°C	5 min	1 cycle	Denaturation	95°C	10 sec	45 cycles	Annealing & Extension	55°C	20 sec
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