

ADPG Pyrophosphorylase Microplate Assay Kit

Catalog # AS0121

Detection and Quantification of ADPG Pyrophosphorylase Activity in Tissue extracts, Cell lysate, Cell culture and Other biological fluids Samples.

This instruction must be read in its entirety before using this product.

For research use only, Not for use in diagnostic procedures.

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I. INTRODUCTION

In enzymology, a glucose-1-phosphate adenylyltransferase (EC 2.7.7.27) is an enzyme that catalyzes the chemical reaction

ATP + alpha-D-glucose 1-phosphate → diphosphate + ADP-glucose

Thus, the two substrates of this enzyme are ATP and alpha-D-glucose 1-phosphate, whereas its two products are diphosphate and ADP-glucose.

This enzyme belongs to the family of transferases, specifically those transferring phosphorus-containing nucleotide groups (nucleotidyltransferases).



II.KIT COMPONENTS

Component	Volume	Storage
96-WellMicroplate	1 plate	
Assay Buffer	30 mlx 4	4 °C
Diluent	20 mlx 1	4 °C
Enzyme	Powder x 1	-20 °C
Substrate	Powder x 1	-20 °C
Technical Manual	1 Manual	

Note:

Enzyme: add 10 ml diluentto dissolve before use.

Substrate: add 10 ml diluentto dissolve before use.

III. MATERIALS REQUIRED BUT NOT PROVIDED

- 1. Microplate reader to read absorbance at 340 nm
- 2. Distilled water
- 3. Pipettor
- 4. Pipette tips
- 5. Mortar
- 6. Centrifuge
- 7. Timer
- 8. Ice



IV. SAMPLE PREPARATION

1.For tissue samples

Weighout 0.1 g tissue, homogenize with 1 mlAssay buffer on ice, centrifuged at 10000g 4°C for 10 minutes, take the supernatant into a new centrifuge tube and keep it on ice for detection.



V. ASSAY PROCEDURE

Add following reagents into the centrifuge tube:

Reagent	Sample		
Sample	50 μl		
Substrate	100 μΙ		
Mix, incubate at 30°C for 30 minutes, put it intoboiling water for 2 minutes. Then			
keep it on ice for cold.Centrifuged at 10000g 4°C for 10 minutes, addthe			
supernatant into the microplate.			
Supernatant	100 μΙ		
Enzyme	100 μΙ		
Mix, measured at 340 nm and recordthe absorbance of 10th second and 130th			
second.			

Note: if the enzyme activity is lower, please add more sample into the reaction system; or increase the reaction time.



VI. CALCULATION

Unit Definition:One Unit of ADPG Pyrophosphorylaseactivity is defined as the enzyme produces 1nmolNADPH per minute.

1. According to the protein concentration of sample

AGPase (U/mg) =
$$(OD_{Sample(130S)} - OD_{Sample(10S)})/(\epsilon \times d) \times V_{Total} \times 10^9/(V_{Sample} \times C_{Protein})/T1/T2 \times 1.5$$

=
$$26.8 \times (OD_{Sample(130S)} - OD_{Sample(10S)}) / C_{Protein}$$

2. According to the weight of sample

AGPase (U/g) =(OD_{Sample(130S)} - OD_{Sample(10S)}) / (
$$\epsilon \times d$$
) $\times V_{Total} \times 10^9$ /(W $\times V_{Sample}$ / V_{Assay})/
T1 / T2 \times 1.5

=
$$26.8 \times (OD_{Sample(130S)} - OD_{Sample(10S)})/W$$

3. According to the quantity of cells or bacteria

AGPase (U/10⁴) =(OD_{Sample(130S)} - OD_{Sample(10S)}) / (
$$\epsilon \times d$$
) $\times V_{Total} \times 10^9$ / (N $\times V_{Sample}$ / V_{Assay})/
T1 / T2 \times 1.5

ε: molar extinction coefficient, 6.22× 10³L/mol/cm;

d: the optical path of 96-Well microplate, 0.6 cm;

C_{Protein}: the protein concentration, mg/ml;

W: the weight of sample, g;

N: the quantity of cell or bacteria, $N \times 10^4$;

V_{Total}: the total volume of the enzymatic reaction, 0.2 ml;

V_{Sample}: the volume of sample, 0.05 ml;

V_{Assay}: the volume of Assay buffer, 1 ml;

T1: the reaction time, 30 minutes.

T2: the reaction time, 2 minutes.



VII. TECHNICAL SUPPORT

For troubleshooting, information or assistance, please go online to www.sabbiotech.cn or contact us at techcn@signalwayantibody.com

VIII. NOTES