Pioneering GTPase and Oncogene Product Development since 2010

Configuration-specific Monoclonal Antibody Based

Arl3 Activation Assay Kit

(30 Assays)

Cat. # 83001

FOR RESEARCH USE ONLY. NOT FOR DIAGNOSTIC APPLICATIONS

NewEast Biosciences
1150 First Avenue, Suite 501
King of Prussia, PA 19406

Support: 610-945-2007
info@neweastbio.com
Web: www.neweastbio.com

$G\alpha_i$ Activation Assay Kit Protocol
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Introduction

A. Background

Arl3 (Arf-like 3) is an ADP-ribosylation factors (Arf) family protein that differs from most Arf family members in the N-terminal extension. Nucleotide exchange of Arl3 is rapid and independent of lipids and detergents. Upon binding of GDT/GTP, Arl3 interacts with and regulates activities of series effector proteins, such as human retinal gene 4 (HRG4), δ-subunit of the cGMP phosphodiesterase (PDEδ), and binder of Arl2 (BART). Arl3 also binds microtubules in a regulated manner to alter specific aspects of cytokinesis via interactions with retinitis pigmentosa 2 (RP2). It has been proposed that RP2 functions in concert with Arl3 to link the cell membrane and the cytoskeleton in photoreceptors as part of the cell signaling or vesicular transport machinery.

Currently there is no direct assay to measure the activation of Arl3 GTPases.

Arl3 Activation Assay Kit is based on the configuration-specific monoclonal antibody that specifically recognizes Arl3-GTP, but not Arl3-GDP. Given the high affinity of monoclonal antibodies to their antigens, the activation assay could be performed in a short time. This assay provides the reliable results with consistent reproducibility.

B. Assay Principle

NewEast Biosciences Arl3 Activation Assay Kit uses configuration-specific anti-Arl3-GTP Mouse monoclonal antibody to measure Arl3-GTP levels in cell extracts or in vitro GTPγS loading Arl3 activation assays. Anti-Arl3-GTP mouse monoclonal antibody is first incubated with cell lysates containing Arl3-GTP. Next, the GTP-bound Arl3 is pulled down by protein A/G agarose. Finally, the precipitated Arl3-GTP is detected through immunoblot analysis using anti-Arl3 mouse monoclonal antibody.

C. Kit Contents

This kit contains enough reagents for approximately 30-35 pull-down assays.

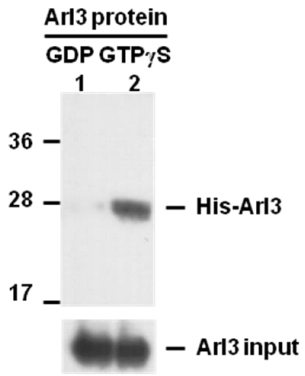
Reagent	Cat. #	Quantity	Storage
Anti-Arl3-GTP Mouse Monoclonal Antibody	26925	30 µL	-20°C
Protein A/G Agarose	30301	600 µL	4°C
5X Assay/Lysis Buffer	30302	30 mL	4°C
Anti-Arl3 Mouse Monoclonal Antibody	26070	50 µL	-20°C
100X GTPγS	30303	50 µL	-20°C
100X GDP	30304	50 µL	-20°C
HRP-Goat Anti-Rabbit IgG	29002	50 µL	-20°C

D. Materials Needed but Not Supplied

1. Stimulated and non-stimulated cell lysates
2. Protease inhibitors
3. 4°C tube rocker or shaker
4. 0.5 M EDTA at pH 8.0
5. 1.0 M $MgCl_2$
6. 2X reducing SDS-PAGE sample buffer
7. Electrophoresis and immunoblotting systems
8. Immunoblotting wash buffer such as TBST (10 mM Tris-HCl, pH 7.4, 0.15 M NaCl, 0.05% Tween-20)
9. Immunoblotting blocking buffer (TBST containing 5% Non-fat Dry Milk or 3% BSA)
10. ECL Detection Reagents

E. Example Results

The following figure demonstrates example results seen with the Arl3 Activation Assay Kit. For reference only.



IP: anti-active Arl3 mAb
IB: anti-Arl3 mAb

Arl3 Activation Assay.
Purified His-tagged Arl3 proteins (Cat. #10152) were immunoprecipitated with the anti-active Arl3 monoclonal antibody (Cat. #26925) after treated with GDP (lane 1) or GTPγS (lane 2) and was blotted with anti-Arl3 monoclonal antibody (Cat. #26070). Input control is shown in bottom panel.

Assay Procedure

A. Reagent Preparation

1X Assay/Lysis Buffer: Shake the 5X Stock Buffer (Cat. # 30302) briefly and dilute with 4 times deionized water to make 1X buffer. Just prior to usage, add protease inhibitors such as 1 mM PMSF, 10 µg/mL leupeptin, and 10 µg/mL aprotinin.

B. Sample Preparation

Adherent Cells

1. Culture cells (one 10-cm plate) to approximately 80-90% confluence (~10^7 cells). Stimulate the cells with activator or inhibitor as desired.
2. Aspirate the culture media and wash twice with ice-cold PBS buffer.
3. Completely remove the final PBS wash and add ice-cold 1X Assay/Lysis Buffer (See Reagent Preparation) to cells (0.5- 1 mL per 10 cm tissue culture plate).
4. Place the culture plates on ice for 10-20 minutes.
5. Detach the cells from the plates by scraping with a cell scraper.
6. Transfer the lysates to appropriate size tubes and place them on ice.
7. If nuclear lysis occurs, the cell lysates may become viscous and difficult to pipette. If this occurs, lysates can be passed through a 27½-gauge syringe needle 3-4 times to shear the genomic DNA.
8. Clear the lysates by centrifuging at 12,000 x g and 4°C for 10 minutes.
9. Collect the supernatant into a tube and store the sample (~1-2 mg of total protein) on ice for immediate use, or snap freeze and store it at -70°C for the future use.

Suspension Cells

1. Culture cells and stimulate with activator or inhibitor as desired.
2. Perform a cell count and then pellet the cells through centrifugation.
3. Aspirate the culture media and wash twice with ice-cold PBS.
4. Completely remove the final PBS wash and add ice-cold 1X Assay/Lysis Buffer (See Reagent Preparation) to the cell pellet (0.5-1 mL per 10^7 cells).
5. Lyse the cells by repeated pipetting.
6. Transfer the lysates to appropriate size tubes and place them on ice.
7. If nuclear lysis occurs, the cell lysates may become viscous and difficult to pipette. If this occurs, lysates can be passed through a 27½-gauge syringe needle 3-4 times to shear the genomic DNA.
8. Clear the lysates by centrifuging at 12,000 x g and 4°C for 10 minutes.
   Collect the supernatant into a tube and store the sample (~1–2 mg of total protein) on ice for immediate use, or snap freeze and store it at -70°C for the future use

C. In vitro GTPγS/GDP Protein for Positive and Negative controls

Note: In vivo stimulation of cells will activate approximately 10% of the available Ar13, whereas in vitro GTPγS protein loading will activate nearly 90% of Ar13.

1. Aliquot 0.5 mL of cell extract (or 1 µg of purified Ar13 protein) into two microcentrifuge tubes.
2. To each tube, add 20 µL of 0.5 M EDTA (final concentration of 20 mM).
3. Add 5 µL of 100 X GTPγS (Cat. # 30303) to the first tube as a positive control.
4. Add 5 µL of 100 X GDP (Cat. # 30304) to the second tube as a negative control.
5. Incubate both tubes at 30°C for 30 minutes with agitation.
6. Stop loading by placing the tubes on ice and adding 32.5 µL of 1 M $MgCl_2$ (final concentration of 60 mM).

D. Affinity Precipitation of Activated G Protein

1. Aliquot 0.5-1 mL of cell lysates (about 1 mg of total cellular protein) to a microcentrifuge tube.
2. Adjust the volume to 1 mL with 1X Assay/Lysis Buffer (See Reagent Preparation).
3. Add 1 µL anti-Ar13-GTP antibody (Cat. # 26925).
4. Prepare the protein A/G Agarose bead slurry (Cat. # 30301) by resuspending through vertexing or titrating.
5. Quickly add 20 µL of resuspended bead slurry to above tube.
6. Incubate the tube at 4°C for 1 hour with gentle agitation.
7. Pellet the beads through centrifugation at 5,000 x g for 1 min.
8. Aspirate and discard the supernatant (making sure not to disturb or remove the bead pellet).
9. Wash the beads 3 times with 0.5 mL of 1X Assay/Lysis Buffer, centrifuging and aspirating each time.
10. After the third wash, pellet the beads through centrifugation and carefully remove all the supernatant.
11. Resuspend the bead pellet in 20 µL of 2X reducing SDS-PAGE sample buffer.
12. Boil the sample for 5 minutes.
13. Centrifuge it at 5,000 x g for 10 seconds.

E. Western Blot Analysis

1. Load 15 µL/well of pull-down supernatant to a polyacrylamide gel (17%). It is recommended to include Prestained Protein Markers (as an indicator of a successful transfer in step 3 below).
2. Perform SDS-PAGE following the manufacturer's instructions.
3. Transfer the gel proteins to a PVDF or nitrocellulose membrane following the manufacturer's instructions.
   Note: Steps 4-11 are at room temperature with agitation
4. Following electroblotting, immerse the PVDF membrane in 100% Methanol for 15 seconds, and then allow it to dry at room temperature for 5 minutes.
   Note: If Nitrocellulose is used instead of PVDF, step 4 Should be skipped.
5. Block the membrane with 5% non-fat dry milk or 3% BSA in TBST for 1 hr at room temperature with constant agitation.
6. Wash the blotted membrane three times with TBST, 5 minutes each time.
7. Incubate the membrane with anti-Ar13 Mouse Monoclonal Antibody (Cat. # 26070), which is freshly diluted 1:50~500 (depending on the amount of Ar13 proteins in your sample) in 5% non-fat dry milk or 3% BSA in TBST, for 1-2 hr at room temperature with constant agitation or at 4°C overnight.
8. Wash the blotted membrane three times with TBST, 5 minutes each time.
9. Incubate the membrane with a secondary antibody (Cat. # 29002), which is freshly diluted 1:1000 in 5% non-fat dry milk or 3% BSA in TBST, for 1 hr at room temperature with constant agitation.
10. Wash the blotted membrane three times with TBST, 5 minutes each time.
11. Use the detection method of your choice such as ECL.