



—A helping hand for your research

Configuration-specific Monoclonal Antibody Based

Sar1 Activation Assay Kit

(30 Assays)

Cat. # 81801

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Sar1 Activation Assay Kit Protocol

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FOR RESEARCH USE ONLY NOT FOR USE IN DIAGNOSTIC PROCEDURES

Product Description

Small GTPases are a superfamily of cellular signaling regulators. Sar1 is a member of these small GTPases. Sar1 functions as a molecular switch to control protein–protein and protein–lipid interactions that direct vesicle budding from the ER. Sar1 is involved in membrane trafficking, and regulates the assembly and disassembly of COPII coats.

Currently there is no direct assay to measure the activation of Sar1 GTPases.

NewEast Biosciences Sar1 Activation Assay Kit is based on the configurationspecific monoclonal antibody that specifically recognizes Sar1–GDP, but not Sar1–GTP. 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.

These anti–Sar1–GTP monoclonal antibodies can also be used to monitor the activation of Sar1 in cells and in tissues by immunohistochemistry.

NewEast Biosciences Sar1 Activation Assay Kit provides a simple and fast method to monitor the activation of Sar1. Each kit provides sufficient quantities to perform 30 assays.

Assay Principle

NewEast Biosciences Sar1 Activation Assay Kit bases on the configurationspecific antiSar1–GDP monoclonal antibody to measure the active Sar1–GTP levels, either from cell extracts or from in vitro GTP γ S loading Sar1 activation assays. Briefly, antiactive Sar1 mouse monoclonal antibody will be incubated with cell lysates containing Sar1–GDP. The bound active Sar1 will then be pulled down by protein A/G agarose. The precipitated active Sar1 will be detected by immunoblot analysis using anti 1ar1 rabbit polyclonal antibody.

Kit Contents

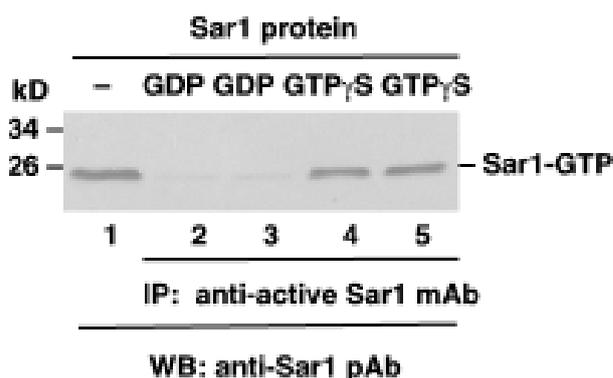
This kit contains enough reagents for approximately 30–35 pull–down assays.

Reagent	Cat. #	Quantity	Storage
Anti–active Sar1 Mouse Monoclonal Antibody	Cat. # 26916	1 X 35 μ l	–20°C
Protein A/G Agarose	Cat. # 30301	1X600 μ l	4°C
5X Assay/Lysis Buffer	Cat. # 30303	1X30mL	4°C
Anti–Sar1 Rabbit polyclonal Antibody	Cat. # 21090	1X50 μ l	–20°C
100x GTP γ S	Cat. # 30302	1X50 μ l	–80°C
100x GDP	Cat. # 30304	1X50 μ l	–80°C
HRP– Goat Anti–Rabbit IgG	Cat. # 29002	1X50 μ l	–20°C

Note: For GDP and GTP γ S, aliquot into 10x5ul volumes, then store at –80 degrees.

Example of Results

The following figure demonstrates typical results seen with NewEast Biosciences Sar1 Activation Assay Kit. One should use the data below for reference only.



Sar1 activation assay. Purified Sar1 proteins were loaded with GDP (lanes 2 and 3) or GTP γ S (lanes 4 and 5).

Immunoprecipitation was done with the antiactive Sar1 monoclonal antibody (Cat. No. 26916). Immunoblot was with an anti1ar1 rabbit polyclonal antibody (Cat. No. 21090).

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Materials Needed but Not Supplied

Stimulated and non-stimulated cell lysates
Protease inhibitors
4°C tube rocker or shaker
0.5 M EDTA, pH8.0
1 M MgCl₂
2X reducing SDS-PAGE sample buffer
Electrophoresis and immunoblotting systems
Immunoblotting wash buffer such as TBST
(10 mM Tris-HCl, pH 7.4, 0.15 M NaCl, 0.05% Tween-20)
Immunoblotting blocking buffer
(TBST containing 5% Non-fat Dry Milk or 3% BSA)
ECL Detection Reagents

A Reagent Preparation

1X Assay/Lysis Buffer: Mix the 5X Stock briefly and dilute to 1X in deionized water. 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, ~ 10⁷ cells) to approximately 80–90% confluence. Stimulate cells with activator or inhibitor as desired.
2. Aspirate the culture media and wash twice with ice-cold PBS.
3. Completely remove the final PBS wash and add ice-cold 1X Assay/Lysis Buffer to the 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 on ice.
7. If nuclear lysis occurs, the cell lysates may become very 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 centrifugation for 10 minutes (12,000 x g at 4°C).
9. Collect the supernatant and store samples (~1–2 mg of total proteins) on ice for immediate use, or snap freeze and store at – 70°C for 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 by 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 to the cell pellet (0.5 – 1 mL per 1 x 10⁷ cells).
5. Lyse the cells by repeated pipetting.
6. Transfer the lysates to appropriate size tubes and place on ice.
7. If nuclear lysis occurs, the cell lysates may become very viscous and difficult to pipette. If this occurs, lysates can be passed through a 27½-gauge syringe needle 3–4 times to shear the
8. Clear the lysates by centrifugation for 10 minutes (12,000 xg at 4°C).
9. Collect the supernatant and store samples on ice for immediate use, or snap freeze and store at – 70°C for future use.

C In vitro GTP γ S/GDP Protein Loading for positive and negative controls

Note: In vivo stimulation of cells will activate approximately

10% of the available Sar1, whereas in vitro GTP γ S protein loading will activate nearly 90% of Sar1

1. Aliquot 0.5 ml of each cell extract to two microfuge tubes (or use 1 µg of purified Sar1 protein).
2. To each tube, add 20 µl of 0.5 M EDTA (to 20 mM final concentration).
3. Add 5 µl of 100 X GTP γ S (to 100 µM, final concentration) to one tube (positive control).
4. Add 5 µl of 100 X GDP (to 1 mM, final concentration) to the second tube (negative control).
5. Incubate the 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 MgCl₂ (to 60 mM, final concentration).

D Affinity Precipitation of Activated G protein

1. Aliquot 0.5 – 1 mL of cell lysate (~1 mg of total cellular protein) to a microcentrifuge tube.
2. Adjust the volume of each sample to 1 mL with 1X Assay/Lysis Buffer.
3. Add 1 µl anti-active Sar1 monoclonal antibody to the tube.
4. Thoroughly resuspend the protein A/G Agarose bead slurry by vortexing or titrating.
5. Quickly add 20 µL of resuspended bead slurry to each tube.
6. Incubate the tubes at 4°C for 1 hour with gentle agitation.
7. Pellet the beads by centrifugation for 1 min at 5,000 x g .
8. Aspirate and discard the supernatant, making sure not to disturb/remove the bead pellet.
9. Wash the bead 3 times with 0.5 mL of 1X Assay/Lysis Buffer, centrifuging and aspirating each time.
10. After the last wash, pellet the beads and carefully remove all the supernatant.
11. Resuspend the bead pellet in 20 µL of 2X reducing SDS-PAGE sample buffer.
12. Boil each sample for 5 minutes.
13. Centrifuge each sample for 10 seconds at 5,000 x g

E Western blot analysis

1. Load 15 µL/well of pull-down supernatant to a polyacrylamide gel (17%). Also, it's recommended to include a pre-stained MW standard (as an indicator of a successful transfer in step 3).2.
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
4. Following the electroblotting step, 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, this step 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. Incubate the membrane with anti- Sar1 polyclonal antibody, freshly diluted 1:50–500 (depending on the amount of Sar1 proteins in your samples) in 5% non-fat dry milk or 3% BSA/TBST, for 1–2 hr at room temperature with constant agitation or at 4°C overnight.
6. Wash the blotted membrane three times with TBST, 5 minutes each time.
7. Incubate the membrane with a secondary antibody (e.g. Goat Anti-Rabbit IgG, HRP-conjugate), freshly diluted 1:1000 in 5% non-fat dry milk or 3% BSA/TBST, for 1 hr at room temperature with constant agitation.
8. Wash the blotted membrane three times with TBST, 5 minutes each time.
9. Use the detection method of your choice such as ECL