

## Anti - p38- $\delta$ (MAPK13)

### Rabbit clonal antibody

#### CAT#

DB 164-0.05 (50  $\mu$ l)  
DB 164-0.1 (100  $\mu$ l)

#### PRODUCT INFORMATION

**Clone number:** I14-L  
**Uniprot:** Human: O15264; Mouse: Q9Z1B7; Rat: Q9WTY9  
**Product description:** Rabbit anti-p38- $\delta$  clonal IgGs  
**Basic information:** Major clone of rabbit immunoglobulin corresponding to immunogenic peptide  
**Immunogen:** Peptide derived from the C-terminus of human p38- $\delta$  sequence. Antibody recognizes the epitope located between Ala353 - Leu365.  
**Species Reactivity:** Human, mouse, rat - tested

**Buffer:** 20 mM Tris-HCl, pH 8.0  
**Stabilizer:** 10 mg/ml BSA  
**Preservative:** 0.05% Sodium Azide

**Storage:** 10  $\mu$ l aliquots at -20°C  
**Handling:** Avoid repeated freezing and thawing  
**Expiration:** 24 months from the shipping date

**Applications:** Western blot, Immunoprecipitation, ELISA, Immunocytochemistry (ICC)

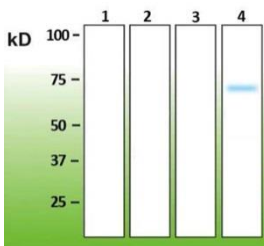
**Dilution range:** Western blotting – 1:1,000  
ELISA – 1:10,000-1:20,000  
Immunoprecipitation – to be tested by user

#### WESTERN BLOT (WB) PROTOCOL - INSTRUCTION MANUAL

##### Western immunoblotting solutions:

- Wash buffer: 1x Tris Buffered Saline (TBS); 0.1% Triton X-100  
- Blocking buffer: 1xTBS; 0.1% Triton X-100; 5% nonfat dry milk

For western blots, incubate the membrane with antibody diluted in blocking buffer for 2 hours at room temperature.

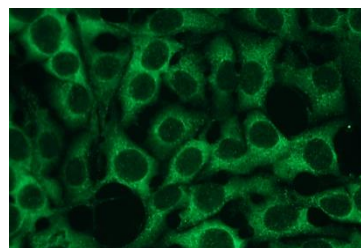


##### Anti - p38- $\delta$ (DB 164)

Western blot analysis of p38- $\delta$ . Lane A – human recombinant p38- $\alpha$  (260-360a.a., N-terminal GST tag; Novus Biologicals, cat. #: H00001432-Q01), lane B – human recombinant p38- $\beta$  (with C-terminal DDK tag, Novus Biologicals, cat. #: NBL1-12870), lane C - human recombinant PKC- $\gamma$  (1-368a.a., N-terminal GST tag, Novus Biologicals, cat. #: H00006300-P01), lane D – human recombinant p38- $\delta$  (1-366a.a., N-terminal GST tag, Novus Biologicals, cat. #: H00005603-P01). 500 ng of recombinant protein was loaded in each well.

#### IMMUNOCYTOCHEMISTRY (ICC) PROTOCOL - INSTRUCTION MANUAL

- Coat coverslips with 1% gelatin-coating solution for 2 hours at room temperature (RT); rinse with distilled water, and let to dry overnight. Before plating the cells, wash the coated coverslips briefly with PBS.
- Fix the cells with 4% paraformaldehyde solution (in PBS, pH 7.2), for 15 min at RT.
- Wash 2 x 3 min with PBS.
- Permeabilize the cells with 0.1% Triton X-100 solution (in PBS, pH 7.2) for 5 min on ice.
- Wash 2 x 3 min with PBS.
- Incubate the cells in blocking buffer (0.3M glycine in PBS, 2% BSA) for 30 min at RT.
- Incubate the cells with primary antibody: anti-p38- $\delta$  clonal antibody at the dilution of **1:200 - 1:500** in antibody dilution buffer (PBS, 1% BSA) for 1 hour at RT in humid chamber.
- Wash 2 x 3 min with PBS.
- Apply the secondary antibody (in this case, the goat anti-rabbit IgG-FITC from Jackson ImmunoResearch, cat. # 111-095-003, was used at 1:300 in antibody dilution buffer, and cells were incubated for 1 hour at RT in dark).
- Wash 3 x 3 min with PBS.
- Rinse once with distilled water.
- Mount the slide for observation, with a drop of anti-fade mounting medium.



Representative picture of p38- $\delta$  expression in HEK293 cells, visualized with clonal rabbit anti-p38- $\delta$  monospecific antibody. Primary antibody dilution - 1:300.

#### PRECAUTIONS

- Intended for professional In Vitro Diagnostic use in laboratories.
- Do not use after expiration date stamped on vial label.
- Avoid contamination of the reagent.
- Any discrepancies in the recommended procedures stated in the working protocol may affect the final results.
- The reagent contains sodium azide ( $\text{NaN}_3$ ) which is highly toxic in higher concentrations. The concentration in the reagent (0.05%) is not considered as hazardous.
- Disposal of waste material must be conducted in accordance with local regulations.
- Wear appropriate Personal Protective Equipment to avoid contact with eyes and skin.