

Recombinant NLRP3 Monoclonal Antibody

catalog number: **AN301382L**

Note: *Centrifuge before opening to ensure complete recovery of vial contents.*

Description

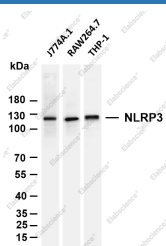
Reactivity	Human;Mouse;Rat
Immunogen	Recombinant Human NLRP3 protein
Host	Rabbit
Isotype	Rabbit IgG
Clone	C111005
Purification	Protein A
Buffer	PBS, 50% glycerol, 0.05% Proclin 300, 0.05% protein protectant.

Applications

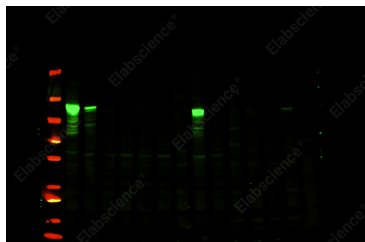
Recommended Dilution

IHC	1:200-1000
WB	1:500-5000
IF	1:200-1000
ELISA	1:5000-20000
IP	1:50-200

Data

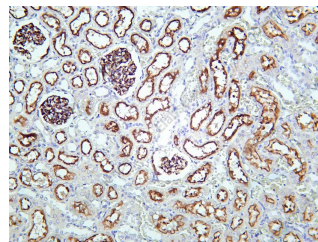


Various whole cell lysates were separated by 4-20% SDS-PAGE, and the membrane was blotted with anti-NLRP3 antibody. The HRP-conjugated Goat anti-Rabbit IgG(H + L) antibody was used to detect the antibody. Lane 1: J774A.1 Lane 2: RAW264.7 Lane 3: THP-1 Predicted band size: 118kDa Observed band size: 118kDa



Western blot analysis of lysate from K562 cells, using NLRP3 Antibody.

Observed-MW:118 kDa
Calculated-MW:118 kDa



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Rev. V1.5

Various whole cell lysates were separated by 4-20% SDS-PAGE, and the primary antibody was used at 4°C, overnight with a 1:2500 dilution. The Dylight 800-conjugated

Goat anti-Rabbit antibody was used to detect the antibody. Lane1: HL60 Lane2: HUVEC Lane3: Human CAF Lane4: HCT116 Lane5: A549 Lane6: Jurkat Lane7: HT22 Lane8: THP-1 Lane9: HK-2 Lane10: PC-12 Lane11: 3T3-L1 Lane12: PC-3 Lane13: U-14 Lane14: 293F. Predicted band size: 118kDa Observed band size: 118kDa

Rat kidney tissue was stained with Anti-NLRP3 rabbit Antibody

Preparation & Storage

Storage Store at -20°C Valid for 12 months. Avoid freeze / thaw cycles.
Shipping Ice bag

Background

This gene encodes a pyrin-like protein containing a pyrin domain, a nucleotide-binding site (NBS) domain, and a leucine-rich repeat (LRR) motif. This protein interacts with the apoptosis-associated speck-like protein PYCARD/ASC, which contains a caspase recruitment domain, and is a member of the NALP3 inflammasome complex. This complex functions as an upstream activator of NF-kappaB signaling, and it plays a role in the regulation of inflammation, the immune response, and apoptosis. Mutations in this gene are associated with familial cold autoinflammatory syndrome (FCAS), Muckle-Wells syndrome (MWS), chronic infantile neurological cutaneous and articular (CINCA) syndrome, and neonatal-onset multisystem inflammatory disease (NOMID). Multiple alternatively spliced transcript variants encoding distinct isoforms have been identified for this gene.