

À des fins de recherche uniquement

# Anticorps Polyclonal de lapin anti-NLRP3

Numéro de catalogue:**CL594-19771** Phare



## Informations de base

|  |                                       |   |
|--|---------------------------------------|---|
| Numéro de catalogue:                           | Numéro d'acquisition GenBank:         | Méthode de purification:                    |
| CL594-19771                                    | NM_001127461                          | Purification par affinité contre l'antigène |
| Taille:  | Identification du gène (NCBI):        | Excitation/Emission maxima wavelengths:     |
| 100ul , Concentration: 1000 µg/ml by Nanodrop; | 114548                                | 588 nm / 604 nm                             |
| Hôte:  | Nom complet:                          |   |
| Lapin  | NLR family, pyrin domain containing 3 |   |
| Isotype:                                       | MW calculé:                           |   |
| IgG  | 118 kDa                               |   |
|  | MW observés:                          |   |
|  | 110 kDa                               |   |

## Applications

Applications testées:

FC (Intra)

Spécificité de l'espèce:

Humain

## Informations générales

NALP3, also named as C1orf7, CIAS1 and PYPAF1, belongs to the NLRP family. NALP3 may function as an inducer of apoptosis. It interacts selectively with ASC and this complex may function as an upstream activator of NF-kappa-B signaling. NALP3 inhibits TNF-alpha induced activation and nuclear translocation of RELA/NF-KB p65. Also inhibits transcriptional activity of RELA. NALP3 activates caspase-1 in response to a number of triggers including bacterial or viral infection which leads to processing and release of IL1B and IL18. Defects in NLRP3 are the cause of familial cold autoinflammatory syndrome type 1 (FCAS1) which also known as familial cold urticaria. Defects in NLRP3 are a cause of Muckle-Wells syndrome (MWS) which is urticaria-deafness-amyloidosis syndrome. Defects in NLRP3 are the cause of chronic infantile neurologic cutaneous and articular syndrome (CINCA) which also known as neonatal onset multisystem inflammatory disease (NOMID). The antibody recognizes the C-term of NALP3.

## Stockage

Stockage:

Stocker à -20 °C. Éviter toute exposition à la lumière. Stable pendant un an après l'expédition.

Tampon de stockage:

PBS avec glycérol à 50 %, Proclin300 à 0,05 % et BSA à 0,5 %, pH 7,3.

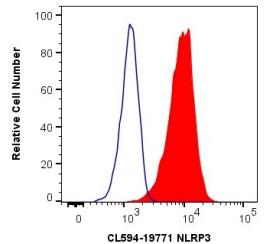
L'aliquotage n'est pas nécessaire pour le stockage à -20°C

\*\*\* Les 20ul contiennent 0,1% de BSA.

For technical support and original validation data for this product please contact:  
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in USA), or 1(312) 455-8498 (outside USA) E: proteintech@ptglab.com  
W: ptglab.com

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## Données de validation sélectionnées



1X $10^6$  THP-1 cells were intracellularly stained with 0.4 ug CoralLite®594 Anti-Human NLRP3 (CL594-19771) (red), or 0.4 ug Isotype Control. Cells were fixed with 4% PFA and permeabilized with Flow Cytometry Perm Buffer (PF00011-C).