For Research Use Only

## SARS-CoV-2 Nucleocapsid Phosphoprotein Recombinant antibody

Catalog Number:80027-1-RR

1 Publications



**Basic Information** 

Catalog Number: 80027-1-RR

Concentration: 1000 ug/ml Source:

Rabbit Isotype:

Immunogen Catalog Number:

AG30676

GenBank Accession Number:

NC\_045512 GeneID (NCBI): 43740575 Full Name:

COVID-19 N Protein

Purification Method:

Protein A purification

CloneNo.: 8C20

Recommended Dilutions:

WB: 1:5000-1:50000 ELISA: 1:10-1:100

**Applications** 

**Tested Applications:** 

WB, ELISA

**Cited Applications:** 

WB

Species Specificity: virus, recombinant protein

Cited Species: human

**Positive Controls:** 

WB: Eukaryotic nucleocapsid phosphoprotein,

Recombinant protein

ELISA: Recombinant protein,

**Background Information** 

The nucleocapsid (N) protein has multiple functions including formation of nucleocapsids, signal transduction virus budding, RNA replication, and mRNA transcription. N protein is an important antigen for coronavirus, and it is normally highly conserved, with a molecular weight of about 50 kDa. it can be used as a marker in diagnostic assays due to its high immunogenicity (PMID: 32416961, PMID: 32235387). A sandwich ELISA for COVID-19 N Protein can be assembled by using 80027-1-RR as capture antibody and conjugated 80026-1-RR for detection.

**Notable Publications** 

Author	Pubmed ID	Journal	Application
Yecheng Zhang	40118151	Virol Sin	WB

Storage

Storage:

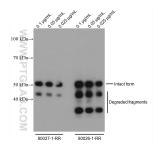
Store at -20°C. Stable for one year after shipment.

Storage Buffer:

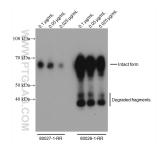
PBS with 0.02% sodium azide and 50% glycerol, pH7.3

Aliquoting is unnecessary for -20°C storage

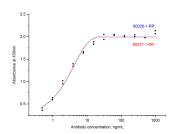
## **Selected Validation Data**



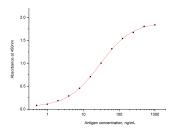
E.coli expressed SARS-CoV-2 Nucleocapsid Phosphoprotein (Cat.NO. Ag30676) was subjected to SDS-PAGE followed by western blot with 80027-1-RR and 80026-1-RR at various work concentration.



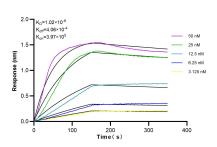
Eukaryotic expressed SARS-CoV-2 Nucleocapsid Phosphoprotein was subjected to SDS-PAGE followed by western blot with 80027-1-RR and 80026-1-RR at various work concentration.



Indirect ELISA was carried out by coating eukaryotic expressed N protein at 70 ng/well followed by blocking and adding serial diluted primary antibody 80026-1-RR and 80027-1-RR respectively. Signal was developed with TMB and stopped by H2SO4. Signal strength was measured by absorbance at 450 nm.



Sandwich ELISA was carried out by coating 80027-1-RR at 80 ng/well followed by blocking and adding different concentration of eukaryotic expressed N protein (0.5-1000 ng/mL). HRP-conjugated80026-1-RR was used at  $1~\mu$  g/mL for detection. Signal was developed with TMB and stopped by H2SO4. Signal strength was measured by absorbance at 450 nm.



Biolayer interferometry (BLL) kinetic assays of 80027-1-RR against SARS-CoV-2 Nucleocapsid Phosphoprotein were performed. The affinity constant is 1.02 nM.