

Recombinant 2019-nCoV N protein

Cat. No. N-127V **Lot. No.** (See product label)

SPECIFICATION

Product Overview	Recombinant 2019-nCoV N protein was expressed in E. coli and purified by Ni column.
Species	Sars-Cov-2
Source	E.coli
Description	<p>Coronaviruses are enveloped viruses with a positive-sense RNA genome and with a nucleocapsid of helical symmetry. Coronavirus nucleoproteins localize to the cytoplasm and the nucleolus, a subnuclear structure, in both virus-infected primary cells and in cells transfected with plasmids that express N protein. Coronavirus N protein is required for coronavirus RNA synthesis, and has RNA chaperone activity that may be involved in template switch. Nucleocapsid protein is a most abundant protein of coronavirus. During virion assembly, N protein binds to viral RNA and leads to formation of the helical nucleocapsid. Nucleocapsid protein is a highly immunogenic phosphoprotein also implicated in viral genome replication and in modulating cell signaling pathways. Because of the conservation of N protein sequence and its strong immunogenicity, the N protein of coronavirus is chosen as a diagnostic tool.</p>
Form	Liquid. PBS, pH 8.0.
Molecular Mass	47.08 kDa
Storage	For short term storage, store at 4 centigrade. For long term storage, store at -20~-70

 Tel: 1-631-559-9269 1-516-512-3133

 Email: info@creative-biomart.com  Fax: 1-631-938-8127

 45-1 Ramsey Road, Shirley, NY 11967, USA

centigrade, avoiding freeze/thaw cycles.

Concentration 1.0 mg/ml

GENE INFORMATION

Gene Name N

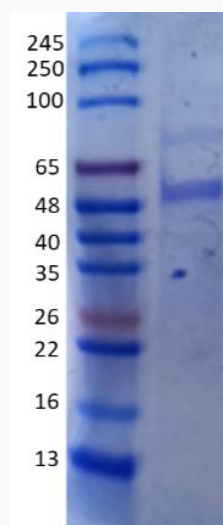
Official Symbol N

Synonyms GU280_gp10

Gene ID 43740575

Protein Refseq YP_009724397.2

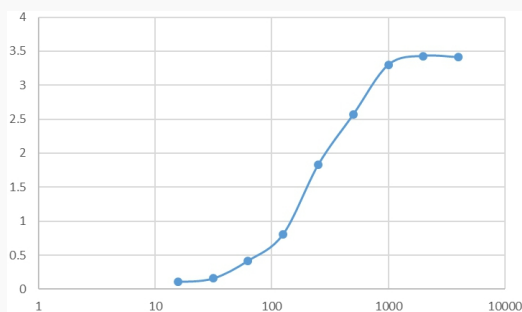
SDS-PAGE of N-127V



 Tel: 1-631-559-9269 1-516-512-3133

 Email: info@creative-biomart.com  Fax: 1-631-938-8127

 45-1 Ramsey Road, Shirley, NY 11967, USA



ELISA of N-127V

ELISA plate was coated by N protein, 100 μ cell, at various concentrations. The direct ELISA analysis was performed by loading 100 μ per well of a anti C-ter His tag HRP conjugated mAb at a concentration of 1:2000. The plate was incubated for 1 hours at 37 centigrade, then washed 5 time. Detection was performed using TMB substrate for 10 minutes at room temperature in the dark. The plate was stopped with 2M sulfuric acid. Absorbances were read on a spectrophotometer at 450 nm.