### For Research Use Only

# G3BP2 Polyclonal antibody

Catalog Number: 16276-1-AP

**Featured Product** 

**15 Publications** 



**Basic Information** 

Catalog Number: GenBank Accession Number: 16276-1-AP BC011731

Size: GeneID (NCBI): 9908

Source: UNIPROT ID: Rabbit Q9UN86

Rabbit Q9UN86

Isotype: Full Name:
IgG GTPase activating protein (SH3

Immunogen Catalog Number:

AG9355

Calculated MW:

482aa,54 kDa; 449aa,51 kDa

domain) binding protein 2

Observed MW: 65-70 kDa

Purification Method:

Antigen affinity purification

Recommended Dilutions:

WB 1:2000-1:16000 IP 0.5-4.0 ug for 1.0-3.0 mg of total

protein lysate IHC 1:250-1:1000 IF 1:50-1:500

**Applications** 

Tested Applications: IF/ICC, IHC, IP, WB, ELISA Cited Applications: WB, IP, IF, RIP, IHC, CoIP Species Specificity: human, mouse

Cited Species: human, mouse

Note-IHC: suggested antigen retrieval with TE buffer pH 9.0; (\*) Alternatively, antigen retrieval may be performed with citrate buffer pH 6.0 Positive Controls:

WB: A549 cells, T47D cells, MCF-7 cells, HeLa cells, Jurkat cells, HEK-293 cells, Neuro-2a cells

IP: HeLa cells.

IHC : human lung cancer tissue, mouse liver tissue, rat

kidney tissue

IF: sodium arsenite treated HeLa cells,

## **Background Information**

Stress granules (SGs) are cytoplasmic mRNA-protein condensates formed in response to cellular stressors, such as oxidative stress, ultraviolet radiation, and viral infection (1). The Ras-GTPase-activating protein-binding proteins (G3BPs), consisting of G3BP1 and G3BP2, are key nucleating factors essential for SG formation. They function to protect RNAs from harmful conditions. G3BP2 is mainly distributed in the cytoplasm and participates in the formation of stress granules, cell differentiation, proliferation, and signal transduction. Accumulating evidence has demonstrated that aberrant expression of G3BP2 contributes to cancer initiation and progression, such as high expression of G3BP2 increasing cell stemness, metastasis and chemoresistance in breast cancer.

#### **Notable Publications**

Author	Pubmed ID	Journal	Application
Hongwei Liu	30458784	Mol Cancer	WB,IHC
Yaqi Chen	35589951	Oncogene	WB,IF,IP
Damin Yun	35782098	PeerJ	WB,IF

Storage

Storage

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

Storage Buffer

PBS with 0.02% sodium azide and 50% glycerol pH 7.3.

Aliquoting is unnecessary for -20°C storage

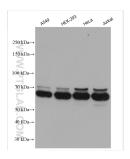
For technical support and original validation data for this product please contact:

T: 4006900926 E: Proteintech-CN@ptglab.com

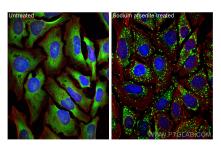
W: ptgcn.coi

This product is exclusively available under Proteintech Group brand and is not available to purchase from any other manufacturer.

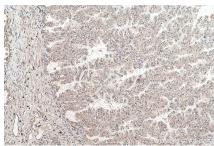
## **Selected Validation Data**



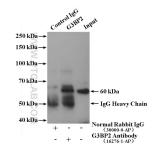
A549 cells were subjected to SDS PAGE followed by western blot with 16276-1-AP (G3BP2 antibody) at dilution of 1:8000 incubated at room temperature for 1.5 burs.



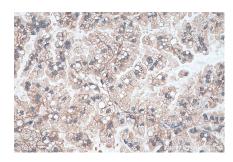
Immunofluorescent analysis of (4% PFA) fixed sodium arsenite treated HeLa cells using G3BP2 antibody (16276-1-AP) at dilution of 1:200 and CoraLite® 488-Conjugated AffiniPure Goat Anti-Rabbit IgG(H+L), CL594-Phalloidin (red).



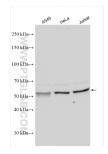
Immunohistochemical analysis of paraffinembedded human lung cancer tissue slide using 16276-1-AP (G3BP2 antibody) at dilution of 1:500 (under 10x lens). Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0).



IP result of anti-G3BP2 (IP:16276-1-AP, 4ug; Detection:16276-1-AP 1:1000) with HeLa cells lysate 2200 ug.



Immunohistochemical analysis of paraffinembedded human lung cancer tissue slide using 16276-1-AP (G3BP2 antibody) at dilution of 1:500 (under 40x lens). Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0).



Various lysates were subjected to SDS PAGE followed by western blot with 16276-1-AP (G3BP2 antibody) at dilution of 1:6000 incubated at room temperature for 1.5 hours.