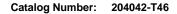
NUP62 Antibody, Rabbit PAb, Antigen Affinity Purified





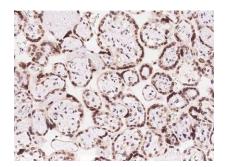
GENERAL INFORMATION	
Immunogen:	E. coli-derived Human NUP62 fragment
Preparation	Produced in rabbits immunized with E. coli-derived Human NUP62 fragment, and purified by antigen affinity chromatography.
Ig Type:	Rabbit IgG
Specificity:	Human NUP62
Formulation:	PBS, pH7.0 with 0.03% Proclin300
Storage:	This antibody can be stored at $2^{\circ}C-8^{\circ}C$ for one month without detectable loss of activity. Antibody products are stable for twelve months from date of receipt when stored at $-20^{\circ}C$ to $-80^{\circ}C$. Preservative-Free. Avoid repeated freeze-thaw cycles.
Alternative Names:	IBSN,p62,SNDI
APPLICATIONS	
Applications:	WB,IHC-P,ICC/IF,IP
RECOMMENDED CONCENTRATION	
IHC-P	IHC-P: 1:50-1:200
ICC/IF	ICC/IF: 1:100-1:500
Western Blot	WB: 1:500-1:2000
Immunoprecipitation	IP:1-5μL/mg of lysate

Please Note: Optimal concentrations/dilutions should be determined by the end user.

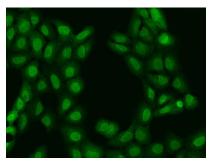
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Catalog Number: 204042-T46

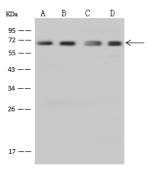




Immunochemical staining of human NUP62 in human placenta with rabbit polyclonal antibody at 1:100 dilution, formalin-fixed paraffin embedded sections.



Immunofluorescence staining of NUP62 in U2OS cells. Cells were fixed with 4% PFA, permeabilzed with 0.1% Triton X-100 in PBS,blocked with 10% serum, and incubated with rabbit anti-Human NUP62 polyclonal antibody (dilution ratio 1:200) at 4°C overnight. Then cells were stained with the Alexa Fluor®488-conjugated Goat Anti-rabbit IgG secondary antibody (green). Positive staining was localized to nuclear membrane.



Anti-NUP62 rabbit polyclonal antibody at 1:500

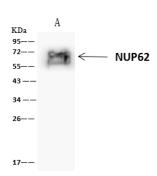
Lane A: A431 Whole Cell Lysate Lane B: HepG2 Whole Cell Lysate Lane C: Jurkat Whole Cell Lysate Lane D: MOLT-4 Whole Cell Lysate

Lysates/proteins at 30 µg per lane.

Secondary Goat Anti-Rabbit IgG (H+L)/HRP at 1/10000 dilution.

Developed using the ECL technique.Performed under reducing conditions.

Predicted band size:53 kDa Observed band size:70 kDa



NUP62 was immunoprecipitated using: Lane A:0.5 mg Jurkat Whole Cell Lysate

4 µL anti-NUP62 rabbit polyclonal antibody and 60 µg of Immunomagnetic beads Protein A/G.Primary antibody:

Anti-NUP62 rabbit polyclonal antibody, at 1:100 dilution

Secondary antibody: Clean-Blot IP Detection Reagent (HRP) at 1:1000dilution

Developed using the ECL technique. Performed under reducing conditions.

Predicted band size: 53 kDa Observed band size: 70 kDa