



NUP62 Monoclonal Antibody

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| Catalog No | BYmab-05869 |
| Isotype | IgG |
| Reactivity | Human;Rat;Mouse |
| Applications | WB |
| Gene Name | NUP62 |
| Protein Name | Nuclear pore glycoprotein p62 (62 kDa nucleoporin) (Nucleoporin Nup62) |
| Immunogen | Synthesized peptide derived from human protein . at AA range: 300-380 |
| Specificity | NUP62 Monoclonal Antibody detects endogenous levels of protein. |
| Formulation | Liquid in PBS containing 50% glycerol, and 0.02% sodium azide. |
| Source | Monoclonal, Mouse,IgG |
| Purification | The antibody was affinity-purified from mouse antiserum by affinity-chromatography using epitope-specific immunogen. |
| Dilution | WB 1:500-2000 |
| Concentration | 1 mg/ml |
| Purity | ≥90% |
| Storage Stability | -20°C/1 year |
| Synonyms | |
| Observed Band | 57kD |
| Cell Pathway | Nucleus, nuclear pore complex . Cytoplasm, cytoskeleton, spindle pole . Nucleus envelope . Cytoplasm, cytoskeleton, microtubule organizing center, centrosome . Central region of the nuclear pore, within the transporter (PubMed:1915414). During mitotic cell division, it associates with the poles of the mitotic spindle (PubMed:24107630). . |
| Tissue Specificity | Brain,Pancreas,Skin,Testis,Urinary bladder, |
| Function | disease:Defects in NUP62 are the cause of infantile striatonigral degeneration (SNDI) [MIM:271930]; also known as infantile bilateral striatal necrosis (IBSN) or infantile bilateral striatal necrosis or familial striatal degeneration. SNDI is a neurological disorder characterized by symmetrical degeneration of the caudate nucleus, putamen, and occasionally the globus pallidus, with little involvement of the rest of the brain. The clinical features include developmental regression, choreoathetosis, dystonia, spasticity, dysphagia, failure to thrive, nystagmus, optic atrophy, and mental retardation.,domain:Contains F-X-F-G repeats.,function:Essential component of the nuclear pore complex. The |

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N-terminal is probably involved in nucleocytoplasmic transport. The C-terminal is probably involved in protein-protein interaction via coiled-coil formation and may function in anchorage of p62 to the

Background

The nuclear pore complex is a massive structure that extends across the nuclear envelope, forming a gateway that regulates the flow of macromolecules between the nucleus and the cytoplasm. Nucleoporins are the main components of the nuclear pore complex in eukaryotic cells. The protein encoded by this gene is a member of the FG-repeat containing nucleoporins and is localized to the nuclear pore central plug. This protein associates with the importin alpha/beta complex which is involved in the import of proteins containing nuclear localization signals. Multiple transcript variants of this gene encode a single protein isoform. [provided by RefSeq, Jul 2008],

matters needing attention

Avoid repeated freezing and thawing!

Usage suggestions

This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.

Products Images