

## TotalSeq™-B0458 anti-Nuclear Pore Complex Proteins Hashtag 8 Antibody

<b>Catalog# / Size</b>	682243 / 10 µg
<b>Clone</b>	MAb414
<b>Regulatory Status</b>	RUO
<b>Other Names</b>	107 kD nucleoporin, NPC proteins, Nuclear pore complex protein Nup107, Nucleoporin 107kD, Nucleoporin Nup107, NUP 107, NUP 84, NUP107, NUP153, NUP155, NUP84, NUP98
<b>Isotype</b>	Mouse IgG1
<b>Barcode Sequence</b>	TGACGCCGTTGTTGT
<b>Description</b>	<p>Nuclear pores are large protein complexes that cross the nuclear envelope. The proteins that make up the nuclear pore complex are known as nucleoporins. About half of the nucleoporins typically contain solenoid protein domains—either an alpha solenoid or a beta-propeller fold, or in some cases both as separate structural domains. Each NPC contains at least 456 individual protein molecules and is composed of 30 distinct proteins (nucleoporins). The other half show structural characteristics typical of "natively unfolded" or intrinsically disordered proteins, i.e. they are highly flexible proteins that lack ordered secondary structure. These disordered proteins are the FG nucleoporins, so called because their amino-acid sequence contains many phenylalanine—glycine repeats.</p> <p>Nuclear pore complexes allow the transport of molecules across the nuclear envelope. This transport includes RNA and ribosomal proteins moving from nucleus to the cytoplasm and proteins (such as DNA polymerase and lamins), carbohydrates, signaling molecules and lipids moving into the nucleus. Although smaller molecules simply diffuse through the pores, larger molecules may be recognized by specific signal sequences and then be diffused with the help of nucleoporins into or out of the nucleus.. Each of the eight protein subunits surrounding the actual pore (the outer ring) projects a spoke-shaped protein over the pore channel.</p> <p>Nucleoporin p62 (p62) protein remains associated with the nuclear pore complex-lamina fraction. p62 is synthesized as a soluble cytoplasmic precursor of 61 kDa followed by modification that involve addition of N-acetylglucosamine residues, followed by association with other complex proteins. 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.</p> <p>P62 is a serine/threonine rich protein of ~520 amino acids, with tetrapeptide repeats on the amino terminus and a series of alpha-helical regions with hydrophobic heptad repeats. P62 assembles into a complex containing 3 additional proteins, p60, p54 and p45 forming the p62 complex of ~235 kDa. Glycosylation appears to be involved in the assembly and disassembly of p62 into higher order complexes, and a serine/threonine rich linker region between Ser270 to Thr294 appear to be regulatory. The p62 complex is localized to both the nucleoplasmic and cytoplasmic sides of the pore complex and the relative diameter of p62 complex relative to the nuclear pore complex suggests it interacts in pore gating.</p> <p>Antibodies to p62 complex are involved in 1 or more autoimmune diseases. P62 glycosylation is increased in diabetes. p62 is also more frequent in Stage IV primary biliary cirrhosis and is prognostic for severe disease. Reduced p62 production has been linked to Alzheimer's disease. It is thought oxidative damage of the p62 promoter is correlated with AD and other neurodegenerative disorders.</p>

### Product Details

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<b>Verified Reactivity</b>	Vertebrate, Xenopus, Yeast
<b>Antibody Type</b>	Monoclonal
<b>Host Species</b>	Mouse
<b>Immunogen</b>	The antibody was raised using a nuclear pore complex mixture.

<b>Formulation</b>	Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide and 1 mM EDTA
<b>Preparation</b>	The antibody was purified by chromatography and conjugated with TotalSeq™-B oligomer under optimal conditions.
<b>Concentration</b>	0.5 mg/mL
<b>Storage &amp; Handling</b>	The antibody solution should be stored undiluted between 2°C and 8°C. <b>Do not freeze.</b>
<b>Application</b>	<a href="#">PG - Quality tested</a>
<b>Recommended Usage</b>	<p>Each lot of this antibody is quality control tested by <a href="#">immunofluorescent staining with flow cytometric analysis</a> and the oligomer sequence is confirmed by sequencing. TotalSeq™-B antibodies are compatible with 10x Genomics Single Cell Gene Expression <a href="#">Solutions</a>.</p> <p>To maximize performance, it is strongly recommended that the reagent be titrated for each application, and that you centrifuge the antibody dilution before adding to the cells at 14,000xg at 2 - 8°C for 10 minutes. Carefully pipette out the liquid avoiding the bottom of the tube and add to the cell suspension. For Proteogenomics analysis, the suggested starting amount of this reagent for titration is ≤ 1.0 µg per million cells in 100 µL volume. Refer to the corresponding TotalSeq™ protocol for specific staining instructions.</p> <p>Buyer is solely responsible for determining whether Buyer has all intellectual property rights that are necessary for Buyer's intended uses of the BioLegend TotalSeq™ products. For example, for any technology platform Buyer uses with TotalSeq™, it is Buyer's sole responsibility to determine whether it has all necessary third party intellectual property rights to use that platform and TotalSeq™ with that platform.</p>
<b>Application Notes</b>	<p>This antibody is effective in immunoblotting (WB), immunohistochemistry (IHC), immunofluorescence (IF), immunoprecipitation (IP) and immunoelectron microscopy (IEM). This antibody has been used successfully with frozen sections.</p> <p>*Predicted MW = 62 kD</p> <p>MAb414 is a reliable general purpose monoclonal antibody which recognizes a related family of NPC proteins. This antibody is ideal for studying the morphology and composition of the nucleus and nuclear envelope. It is also useful in studying changes in the nuclear structure during mitosis and meiosis.</p>
<b>Additional Product Notes</b>	<p>TotalSeq™ reagents are designed to profile protein levels at a single cell level following an optimized protocol similar to the CITE-seq workflow. A compatible single cell device (e.g. <a href="#">10x Genomics Chromium System and Reagents</a>) and sequencer (e.g. Illumina analyzers) are required. Please contact <a href="#">technical support</a> for more information, or visit <a href="#">biolegend.com/totalseq</a>.</p> <p>The barcode flanking sequences are GTGACTGGAGTTCAGACGTGTGCTCTTCCGATCTNNNNNNNNNN (PCR handle), and NNNNNNNNGCTTTAAGGCCGTCCTAGC*A*A (capture sequence). N represents either randomly selected A, C, G, or T, and * indicates a phosphorothioated bond, to prevent nuclease degradation.</p> <p>View more applications data for this product in our <a href="#">Scientific Poster Library</a>.</p>
<b>Application References</b>	<ol style="list-style-type: none"> <li>Zheng X, <i>et al.</i> 2012. <i>J. Biol. Chem.</i> 287:38254. (IF) <a href="#">PubMed</a></li> <li>Kimura T, <i>et al.</i> 2003. <i>Mol Cell Biol.</i> 23:1304. (IHC) <a href="#">PubMed</a></li> <li>Lopez-Soler RI, <i>et al.</i> 2001. <i>J Cell Biol.</i> 154:61. (IF, WB) <a href="#">PubMed</a></li> <li>Aris JP, Blobel G. 1989. <i>J Cell Biol.</i> 108:2059. (WB, IP, IF, EM) <a href="#">PubMed</a></li> <li>Davis LI, Blobel G. 1987. <i>PNAS USA.</i> 84:7552. (IP, IF) <a href="#">PubMed</a></li> <li>Edens LJ, Levy DL. 2014. <i>J. Cell. Biol.</i> 206:473.</li> <li>Davis LI, Blobel G. 1986. <i>Cell.</i> 45:699.</li> <li>Blobel G. 1985. <i>PNAS USA</i> 82:8527.</li> </ol>
<b>RRID</b>	AB_2892488 (BioLegend Cat. No. 682243)

## Antigen Details

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<b>Biology Area</b>	Cell Biology, Cell Motility/Cytoskeleton/Structure, Neuroscience, Neuroscience Cell Markers
<b>Molecular Family</b>	Nuclear Markers
<b>Antigen References</b>	<ol style="list-style-type: none"> <li>Yoshimura S, <i>et al.</i> 2013. <i>J. Cell Sci.</i> 126:3141-50.</li> </ol>
<b>Gene ID</b>	NA

## Related Protocols

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[TotalSeq™-B or -C with 10x Feature Barcoding Technology](#)

## Other Formats

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Purified anti-Nuclear Pore Complex Proteins , Alexa Fluor® 594 anti-Nuclear Pore Complex Proteins , Direct-Blot™ HRP anti-Nuclear Pore Complex Proteins , Alexa Fluor® 647 anti-Nuclear Pore Complex Proteins, TotalSeq™-A0451 anti-Nuclear Pore Complex Proteins Hashtag 1, TotalSeq™-A0452 anti-Nuclear Pore Complex Proteins Hashtag 2, TotalSeq™-A0453 anti-Nuclear Pore Complex Proteins Hashtag 3, TotalSeq™-A0454 anti-Nuclear Pore Complex Proteins Hashtag 4, TotalSeq™-A0456 anti-Nuclear Pore Complex Proteins Hashtag 6, TotalSeq™-A0457 anti-Nuclear Pore Complex Proteins Hashtag 7, TotalSeq™-A0458 anti-Nuclear Pore Complex Proteins Hashtag 8, TotalSeq™-A0459 anti-Nuclear Pore Complex Proteins Hashtag 9, TotalSeq™-A0460 anti-Nuclear Pore Complex Proteins Hashtag 10, TotalSeq™-A0455 anti-Nuclear Pore Complex Proteins Hashtag 5, TotalSeq™-A0462 anti-Nuclear Pore Complex Proteins Hashtag 12, TotalSeq™-A0463 anti-Nuclear Pore Complex Proteins Hashtag 13, TotalSeq™-A0465 anti-Nuclear Pore Complex Proteins Hashtag 15, TotalSeq™-A0461 anti-Nuclear Pore Complex Proteins Hashtag 11, TotalSeq™-A0464 anti-Nuclear Pore Complex Proteins Hashtag 14, TotalSeq™-B0453 anti-Nuclear Pore Complex Proteins Hashtag 3, TotalSeq™-B0452 anti-Nuclear Pore Complex Proteins Hashtag 2, TotalSeq™-B0451 anti-Nuclear Pore Complex Proteins Hashtag 1, TotalSeq™-B0458 anti-Nuclear Pore Complex Proteins Hashtag 8, TotalSeq™-B0457 anti-Nuclear Pore Complex Proteins Hashtag 7

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