

#68591 Store at -20°C

Lamin B1 (119D5-F1) Mouse mAb



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| Applications: IF-IC | Reactivity: H | Sensitivity: Endogenous | Source/Isotype: Mouse IgG1 | UniProt ID: #P20700 | Entrez-Gene Id: 4001 |
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| Product Usage Information | Application Immunofluorescence (Immunocytochemistry) | Dilution 1:400 - 1:1600 |
| Storage | Supplied at 1 mg/mL in PBS containing 0.09% sodium azide. Store at -20°C. <i>This product will freeze at -20°C so it is recommended to aliquot into single-use vials to avoid multiple freeze/thaw cycles.</i> A slight precipitate may be present, but will not interfere with antibody performance. This product is stable for 36 months when stored at -20°C. | |
| Specificity / Sensitivity | Lamin B1 (119D5-F1) Mouse mAb recognizes endogenous levels of total Lamin B1 by immunofluorescence. Testing by immunofluorescence was validated for human cells. Reactivity is expected in other species, including mouse and rat. | |
| Source / Purification | Monoclonal antibody is produced by immunizing animals with purified rat liver lamins. | |
| Background | Lamins are nuclear membrane structural components that are important in maintaining normal cell functions, such as cell cycle control, DNA replication, and chromatin organization (1-3). Lamins have been subdivided into types A and B. Type-A lamins consist of lamin A and C, which arise from alternative splicing of the lamin A gene <i>LMNA</i> . Lamin A and C are cleaved by caspases into large (41-50 kDa) and small (28 kDa) fragments, which can be used as markers for apoptosis (4,5). Type-B lamins consist of lamin B1 and B2, encoded by separate genes (6-8). Lamin B1 is also cleaved by caspases during apoptosis (9). Research studies have shown that duplication of the lamin B1 gene <i>LMNB1</i> is correlated with pathogenesis of the neurological disorder adult-onset leukodystrophy (10). | |
| Background References | <ol style="list-style-type: none"> Gruenbaum, Y. et al. (2000) <i>J Struct Biol</i> 129, 313-23. Goldberg, M. et al. (1999) <i>Crit Rev Eukaryot Gene Expr</i> 9, 285-93. Yabuki, M. et al. (1999) <i>Physiol Chem Phys Med NMR</i> 31, 77-84. Rao, L. et al. (1996) <i>J Cell Biol</i> 135, 1441-55. Orth, K. et al. (1996) <i>J Biol Chem</i> 271, 16443-6. Biamonti, G. et al. (1992) <i>Mol Cell Biol</i> 12, 3499-506. Lin, F. and Worman, H.J. (1995) <i>Genomics</i> 27, 230-6. Pollard, K.M. et al. (1990) <i>Mol Cell Biol</i> 10, 2164-75. Chandler, J.M. et al. (1997) <i>Biochem J</i> 322 (Pt 1), 19-23. Padiath, Q.S. et al. (2006) <i>Nat Genet</i> 38, 1114-23. | |

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| Species Reactivity | Species reactivity is determined by testing in at least one approved application (e.g., western blot). |
| Applications Key | IF-IC: Immunofluorescence (Immunocytochemistry) |
| Cross-Reactivity Key | H: human M: mouse R: rat Hm: hamster Mk: monkey Vir: virus Mi: mink C: chicken Dm: D. melanogaster X: Xenopus Z: zebrafish B: bovine Dg: dog Pg: pig Sc: S. cerevisiae Ce: C. elegans Hr: horse GP: Guinea Pig Rab: rabbit All: all species expected |
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