Lamin B1 Rabbit pAb

Catalog No.: A1910 40 Publications



Basic Information

Observed MW

68kDa/45kDa

Calculated MW

66kDa

Category

Primary antibody

Applications

ELISA, WB, IHC-P, IF/ICC, IP, ChIP

Cross-Reactivity

Human, Mouse, Rat

Background

This gene encodes one of the two B-type lamin proteins and is a component of the nuclear lamina. A duplication of this gene is associated with autosomal dominant adult-onset leukodystrophy (ADLD). Alternative splicing results in multiple transcript variants.

Recommended Dilutions

WB	1:500 - 1:1000

IHC-P 1:50 - 1:200

IF/ICC 1:50 - 1:200

IP 0.5μg-4μg antibody for

200μg-400μg extracts of

whole cells

ChIP 5μg antibody for

10μg-15μg of Chromatin

Immunogen Information

Gene ID	Swiss Prot
4001	P20700

Immunogen

Recombinant fusion protein containing a sequence corresponding to amino acids 397-586 of human Lamin B1 (NP_005564.1).

Synonyms

LMN; ADLD; LMN2; LMNB; MCPH26; Lamin B1

Contact

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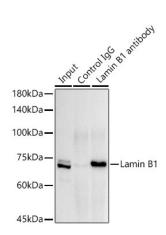
Product Information

SourceIsotypePurificationRabbitIgGAffinity purification

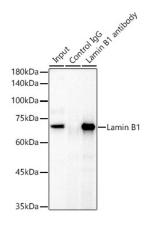
Storage

Store at -20°C. Avoid freeze / thaw cycles.

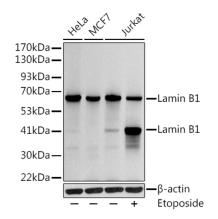
Buffer: PBS with 0.01% thimerosal,50% glycerol,pH7.3.



Immunoprecipitation analysis of 300 μ g extracts of HeLa cells using 3 μ g Lamin B1 antibody (A1910). Western blot was performed from the immunoprecipitate using Lamin B1 (A1910) at a dilution of 1:1000.



Immunoprecipitation analysis of 300 µg extracts of HeLa cells using 3 µg Lamin B1 antibody (A1910). Western blot was performed from the immunoprecipitate using Lamin B1 antibody (A1910) at a dilution of 1.1000



Western blot analysis of various lysates using Lamin B1 Rabbit pAb (A1910) at 1:1000 dilution. Jurkat cells were treated by Etoposide (25 uM) at 37° C for 5 hours.

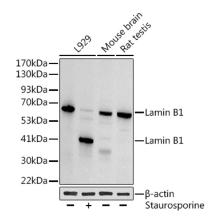
Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

Lysates/proteins: 25µg per lane.

Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

Exposure time: 10s.



Western blot analysis of various lysates using Lamin B1 Rabbit pAb (A1910) at 1:1000 dilution.L929 cells were treated by staurosporine(1 uM) for 3 hour.

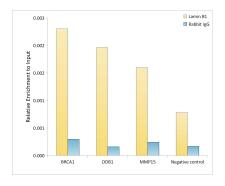
Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

Lysates/proteins: 25µg per lane.

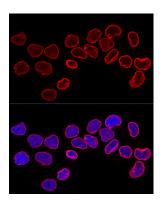
Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

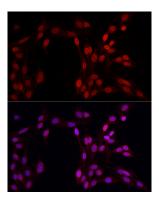
Exposure time: 30s.



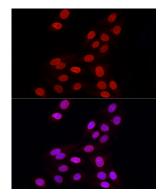
Chromatin immunoprecipitation analysis of extracts of HeLa cells, using Lamin B1 Rabbit pAb antibody (A1910) and rabbit IgG.The amount of immunoprecipitated DNA was checked by quantitative PCR. Histogram was constructed by the ratios of the immunoprecipitated DNA to the input.



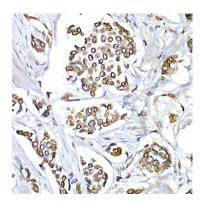
Confocal immunofluorescence analysis of HeLa cells using Lamin B1 Rabbit pAb (A1910) at dilution of 1:200. Blue: DAPI for nuclear staining.



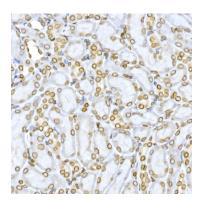
Immunofluorescence analysis of PC-12 cells using Lamin B1 Rabbit pAb (A1910) at dilution of 1:100 (40x lens). Secondary antibody: Cy3 Goat Anti-Rabbit IgG (H+L) (AS007) at 1:500 dilution. Blue: DAPI for nuclear staining.



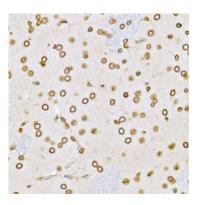
Immunofluorescence analysis of U2OS cells using Lamin B1 Rabbit pAb (A1910) at dilution of 1:100 (40x lens). Secondary antibody: Cy3 Goat Anti-Rabbit IgG (H+L) (AS007) at 1:500 dilution. Blue: DAPI for nuclear staining.



Immunohistochemistry analysis of Lamin B1 in paraffin-embedded human breast cancer using Lamin B1 Rabbit pAb (A1910) at dilution of 1:150 (40x lens).Perform high pressure antigen retrieval with 10 mM citrate buffer pH 6.0 before commencing with IHC staining protocol.



Immunohistochemistry analysis of Lamin B1 in paraffin-embedded mouse kidney using Lamin B1 Rabbit pAb (A1910) at dilution of 1:150 (40x lens).Perform high pressure antigen retrieval with 10 mM citrate buffer pH 6.0 before commencing with IHC staining protocol.



Immunohistochemistry analysis of Lamin B1 in paraffin-embedded rat brain using Lamin B1 Rabbit pAb (A1910) at dilution of 1:150 (40x lens).Perform high pressure antigen retrieval with 10 mM citrate buffer pH 6.0 before commencing with IHC staining protocol