# Lamin B2 Rabbit mAb

Catalog No.: A5001 Recombinant 1 Publications



## **Basic Information**

### **Observed MW**

68kDa

### **Calculated MW**

70kDa

### Category

Primary antibody

### **Applications**

ELISA,WB,IHC-P,IF/ICC

### **Cross-Reactivity**

Human, Mouse, Rat

#### CloneNo number

ARC1252

## **Background**

This gene encodes a B type nuclear lamin. The nuclear lamina consists of a two-dimensional matrix of proteins located next to the inner nuclear membrane. The lamin family of proteins make up the matrix and are highly conserved in evolution. During mitosis, the lamina matrix is reversibly disassembled as the lamin proteins are phosphorylated. Lamin proteins are thought to be involved in nuclear stability, chromatin structure and gene expression. Vertebrate lamins consist of two types, A and B. Mutations in this gene are associated with acquired partial lipodystrophy.

## **Recommended Dilutions**

WB	1:500 - 1:1000	
IHC-P	1:50 - 1:200	
IF/ICC	1:50 - 1:200	

## **Immunogen Information**

Gene ID	Swiss Prot
84823	003252

### **Immunogen**

A synthetic peptide corresponding to a sequence within amino acids 521-620 of human Lamin B2 (Q03252).

## **Synonyms**

EPM9; LMN2; LAMB2; MCPH27; Lamin B2

## **Contact**

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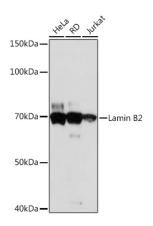
### **Product Information**

Source	Isotype	Purification
Rabbit	IgG	Affinity purification

#### Storage

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

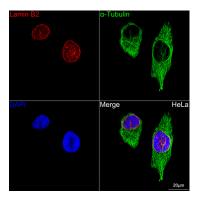
Buffer: PBS with 0.02% sodium azide, 0.05% BSA, 50% glycerol, pH7.3.



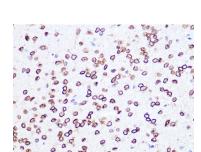
Western blot analysis of various lysates using Lamin B2 Rabbit mAb (A5001) at 1:1000 dilution. Secondary antibody: HRP Goat Anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

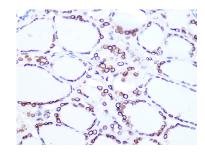
Lysates/proteins: 25µg per lane.

Exposure time: 30s.



Blocking buffer: 3% nonfat dry milk in TBST. Detection: ECL Basic Kit (RM00020).

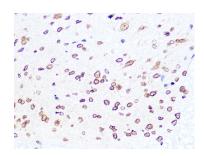


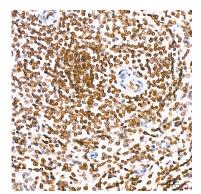


Confocal imaging of HeLa cells using Lamin B2 Rabbit mAb (A5001, dilution 1:100) (Red). The cells were counterstained with α-Tubulin Rabbit mAb (AC049, dilution 1:100) (Green). DAPI was used for nuclear staining (blue). Objective: 60x.

Immunohistochemistry analysis of paraffinembedded Rat brain using Lamin B2 Rabbit mAb (A5001) at dilution of 1:100 (40x lens).Perform microwave antigen retrieval with 10 mM PBS buffer pH 7.2 before commencing with IHC staining protocol.

Immunohistochemistry analysis of paraffinembedded Human thyroid cancer using Lamin B2 Rabbit mAb (A5001) at dilution of 1:100 (40x lens).Perform microwave antigen retrieval with 10 mM PBS buffer pH 7.2 before commencing with IHC staining protocol.

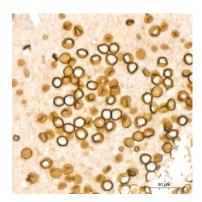




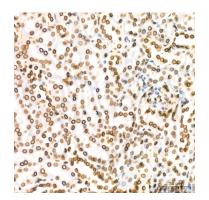
Immunohistochemistry analysis of paraffinembedded Mouse spinal cord using Lamin B2 Rabbit mAb (A5001) at dilution of 1:100 (40x lens).Perform microwave antigen retrieval with 10 mM PBS buffer pH 7.2 before commencing with IHC staining protocol.

Immunohistochemistry analysis of paraffinembedded Human colon carcinoma tissue using Lamin B2 Rabbit mAb (A22347) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.

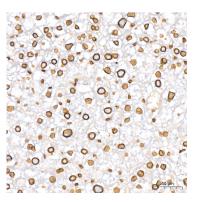
Immunohistochemistry analysis of paraffinembedded Human spleen tissue using Lamin B2 Rabbit mAb (A22347) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.



Immunohistochemistry analysis of paraffinembedded Mouse brain tissue using Lamin B2 Rabbit mAb (A22347) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.



Immunohistochemistry analysis of paraffinembedded Mouse kidney tissue using Lamin B2 Rabbit mAb (A22347) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.



Immunohistochemistry analysis of paraffinembedded Mouse liver tissue using Lamin B2 Rabbit mAb (A22347) at a dilution of 1:200 (40x lens). High pressure antigen retrieval was performed with 0.01 M citrate buffer (pH 6.0) prior to IHC staining.