

Lamin A/C (5D12) Mouse mAb
Catalog # AP53523**Specification**

Lamin A/C (5D12) Mouse mAb - Product Information

Application	WB, IF
Reactivity	Rat
Host	Mouse
Clonality	Monoclonal Antibody

Lamin A/C (5D12) Mouse mAb - Additional Information**Other Names**

FPL; IDC; LFP; CDDC; EMD2; FPLD; HGPS; LDP1; LMN1; LMNC; PRO1; CDCD1; CMD1A; FPLD2; LMNL1; CMT2B1; LGMD1B

Dilution

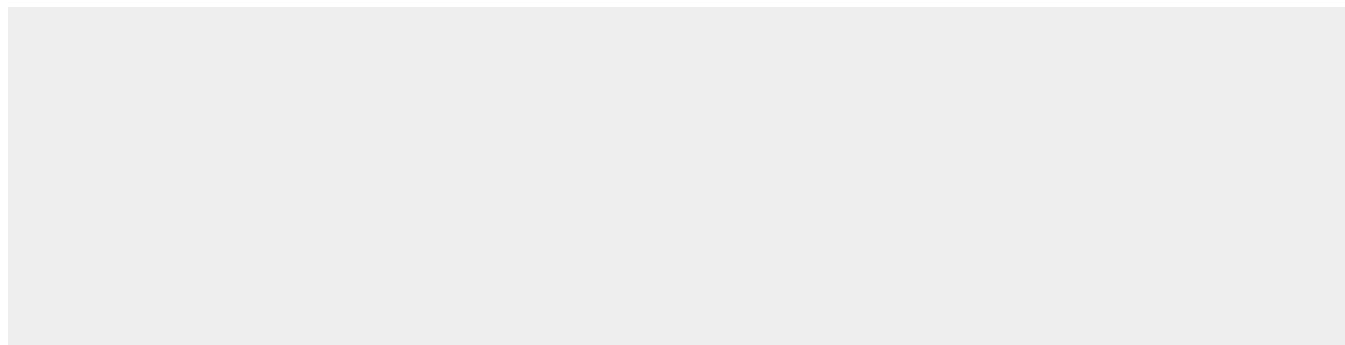
WB~~1:1000

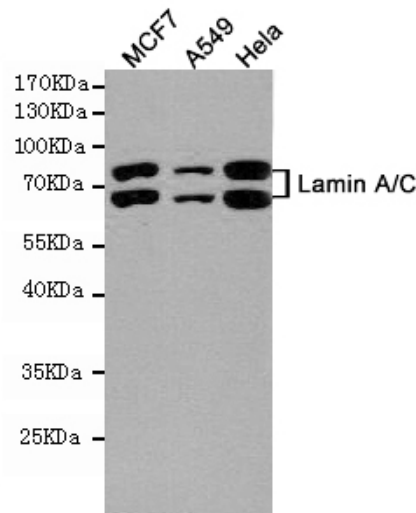
IF~~1:200

Lamin A/C (5D12) Mouse mAb - Protein Information**Lamin A/C (5D12) Mouse mAb - Protocols**

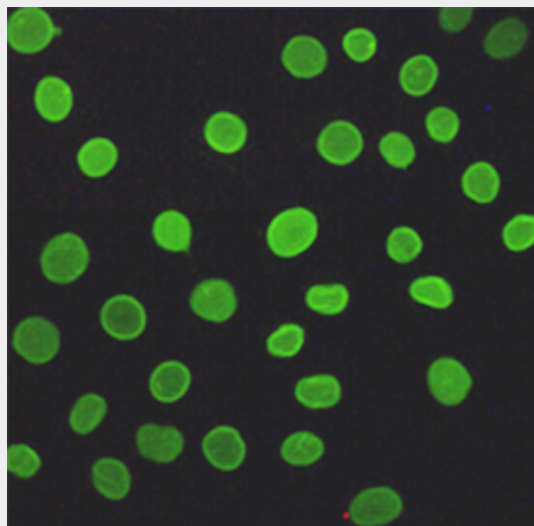
Provided below are standard protocols that you may find useful for product applications.

- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

Lamin A/C (5D12) Mouse mAb - Images



Western blot detection of Lamin A/C in MCF7, A549 and HeLa cell lysates using Lamin A/C mouse mAb (1:1000 diluted). Predicted band size: 74,63KDa. Observed band size: 74,63KDa.



Immunofluorescent analysis of A549 cells fixed with 4% Paraformaldehyde and using anti-Lamin A/C mouse mAb (dilution 1:200).

Lamin A/C (5D12) Mouse mAb - Background

Lamins are components of the nuclear lamina, a fibrous layer on the nucleoplasmic side of the inner nuclear membrane, which is thought to provide a framework for the nuclear envelope and may also interact with chromatin. Lamin A and C are present in equal amounts in the lamina of mammals. Plays an important role in nuclear assembly, chromatin organization, nuclear membrane and telomere dynamics. Required for normal development of peripheral nervous system and skeletal muscle and for muscle satellite cell proliferation. Required for osteoblastogenesis and bone formation. Also prevents fat infiltration of muscle and bone marrow, helping to maintain the volume and strength of skeletal muscle and bone. Prelamin-A/C can accelerate smooth muscle cell senescence. It acts to disrupt mitosis and induce DNA damage in vascular smooth muscle cells (VSMCs), leading to mitotic failure, genomic instability, and premature senescence.