

ATRX monoclonal antibody, clone CL0537

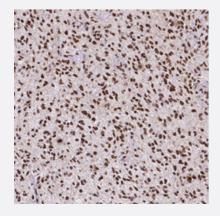
Catalog # MAB15638 Size 100 uL

Applications



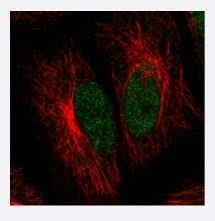
Western Blot (Cell lysate)

Western Blot analysis of A-549 cell lysate with ATRX monoclonal antibody, clone CL0537 (Cat # MAB15638).



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections)

Immunohistochemical staining (Formalin-fixed paraffin-embedded sections) of human glioma with ATRX monoclonal antibody, clone CL0537 (Cat # MAB15638) shows strong nuclear positivity in tumor cells.



Immunofluorescence

Immunofluorescent staining of HeLa cells with ATRX monoclonal antibody, clone CL0537 (Cat # MAB15638) (Green) shows clear nuclear (without nucleoli) staining. Microtubule and nuclear probes are visualized in red and blue respectively (where available).

Specification



Product Information

Product Description	Mouse monoclonal antibody raised against partial recombinant human ATRX.
Immunogen	Recombinant protein corresponding to human ATRX.
Sequence	AAWAEYEAEKKGLTMRFNIPTGTNLPPVSFNSQTPYIPFNLGALSAMSNQQLEDLINQGREKVVEA TNSVTAVRIQPLEDIISAVWKENMNLSEAQVQALALSRQASQELDVKRREAIYNDVLTKQQMLISC VQRILMNRR
Host	Mouse
Reactivity	Human
Form	Liquid
Purification	Protein A purification
Isotype	lgG1
Recommend Usage	Immunofluorescence (1-4 ug/mL) Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) (1:200-1:500) Western Blot (1:500-1:1000) The optimal working dilution should be determined by the end user.
Storage Buffer	In PBS, pH 7.2 (40% glycerol, 0.02% sodium azide).
Storage Instruction	Store at 4°C. For long term storage store at -20°C. Aliquot to avoid repeated freezing and thawing.
Note	This product contains sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which shoul d be handled by trained staff only.

Applications

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Gene Info — ATRX	
Entrez GenelD	<u>546</u>
Protein Accession#	P46100
Gene Name	ATRX
Gene Alias	ATR2, MGC2094, MRXHF1, RAD54, RAD54L, SFM1, SHS, XH2, XNP, ZNF-HX
Gene Description	alpha thalassemia/mental retardation syndrome X-linked (RAD54 homolog, S. cerevisiae)
Omim ID	300032 300448 301040 309580
Gene Ontology	Hyperlink
Gene Summary	The protein encoded by this gene contains an ATPase/helicase domain, and thus it belongs to the SWI/SNF family of chromatin remodeling proteins. The mutations of this gene are associated with an X-linked mental retardation (XLMR) syndrome most often accompanied by alpha-thalassemia (ATRX) syndrome. These mutations have been shown to cause diverse changes in the pattern of DNA methylation, which may provide a link between chromatin remodeling, DNA methylation, and gene expression in developmental processes. This protein is found to undergo cell cycle-dependent phosphorylation, which regulates its nuclear matrix and chromatin association, and suggests its involvement in the gene regulation at interphase and chromosomal segregation in mitosis. Multiple alternatively spliced transcript variants encoding distinct isoforms have been reported. [provided by RefSeq
Other Designations	DNA dependent ATPase and helicase OTTHUMP00000024265 OTTHUMP00000062079 X-link ed nuclear protein Zinc finger helicase helicase 2, X-linked transcriptional regulator ATRX

Disease

- Breast cancer
- Breast Neoplasms
- Genetic Predisposition to Disease