

Recombinant Anti-LOXL2 Rabbit mAb

Catalog # AP94855

Product Information

Application	WB, IHC, FC, IF/IC
Primary Accession	Q9Y4K0
Reactivity	Human, Mouse, Rat
Host	Rabbit
Clonality	Monoclonal
Calculated MW	86725

Additional Information

Gene ID	4017
Other Names	Lysyl oxidase homolog 2; Lysyl oxidase-like protein 2; Lysyl oxidase-related protein 2; Lysyl oxidase-related protein WS9-14
Target/Specificity	KLH-conjugated synthetic peptide encompassing a sequence within human LOXL2 protein. The exact sequence is proprietary.
Dilution	WB~~WB (1/500 - 1/1000) IHC~~1:100~500 FC~~1:10~50 IF/IC~~N/A
Format	Liquid in PBS, pH 7.4, containing 50% glycerol, 0.2% BSA and 0.01% sodium azide.
Storage	Store at -20 °C.Stable for 12 months from date of receipt

Protein Information

Name	LOXL2
Function	Mediates the post-translational oxidative deamination of lysine residues on target proteins leading to the formation of deaminated lysine (allysine) (PubMed:27735137). Acts as a transcription corepressor and specifically mediates deamination of trimethylated 'Lys-4' of histone H3 (H3K4me3), a specific tag for epigenetic transcriptional activation (PubMed:27735137). Shows no activity against histone H3 when it is trimethylated on 'Lys-9' (H3K9me3) or 'Lys-27' (H3K27me3) or when 'Lys-4' is monomethylated (H3K4me1) or dimethylated (H3K4me2) (PubMed:27735137). Also mediates deamination of methylated TAF10, a member of the transcription factor IID (TFIID) complex, which induces release of TAF10 from promoters, leading to inhibition of TFIID-dependent transcription (PubMed:25959397). LOXL2-mediated deamination of TAF10 results in transcriptional repression of genes required for embryonic stem cell pluripotency including POU5F1/OCT4, NANOG, KLF4 and SOX2 (By similarity). Involved in epithelial to mesenchymal transition (EMT) via interaction with SNAI1 and participates in repression of

E-cadherin CDH1, probably by mediating deamination of histone H3 (PubMed:[16096638](#), PubMed:[24414204](#), PubMed:[27735137](#)). During EMT, involved with SNAI1 in negatively regulating pericentromeric heterochromatin transcription (PubMed:[24239292](#)). SNAI1 recruits LOXL2 to pericentromeric regions to oxidize histone H3 and repress transcription which leads to release of heterochromatin component CBX5/HP1A, enabling chromatin reorganization and acquisition of mesenchymal traits (PubMed:[24239292](#)). Interacts with the endoplasmic reticulum protein HSPA5 which activates the IRE1-XBP1 pathway of the unfolded protein response, leading to expression of several transcription factors involved in EMT and subsequent EMT induction (PubMed:[28332555](#)). Involved in E-cadherin repression following hypoxia, a hallmark of EMT believed to amplify tumor aggressiveness, suggesting that it may play a role in tumor progression (PubMed:[20026874](#)). When secreted into the extracellular matrix, promotes cross-linking of extracellular matrix proteins by mediating oxidative deamination of peptidyl lysine residues in precursors to fibrous collagen and elastin (PubMed:[20306300](#)). Acts as a regulator of sprouting angiogenesis, probably via collagen IV scaffolding (PubMed:[21835952](#)). Acts as a regulator of chondrocyte differentiation, probably by regulating expression of factors that control chondrocyte differentiation (By similarity).

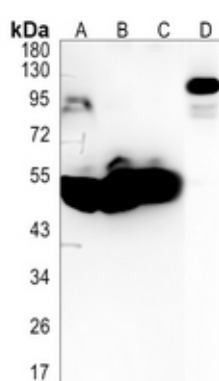
Cellular Location

Secreted, extracellular space, extracellular matrix, basement membrane. Nucleus. Chromosome. Endoplasmic reticulum. Note=Associated with chromatin (PubMed:[27735137](#)). It is unclear how LOXL2 is nuclear as it contains a signal sequence and has been shown to be secreted (PubMed:[23319596](#)) However, a number of reports confirm its intracellular location and its key role in transcription regulation (PubMed:[22204712](#), PubMed:[22483618](#)).

Tissue Location

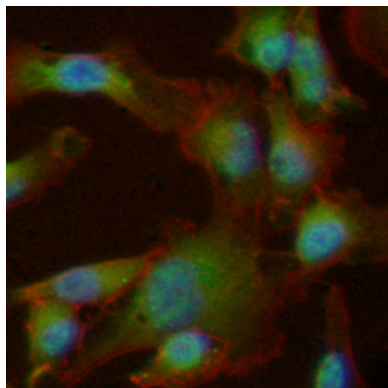
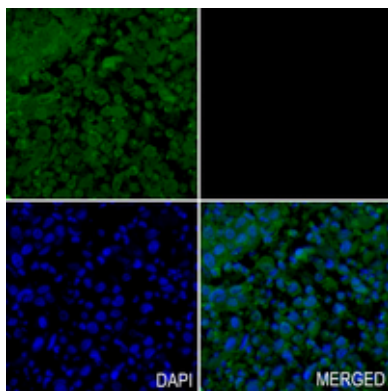
Expressed in many tissues (PubMed:[10212285](#)). Highest expression in reproductive tissues, placenta, uterus and prostate (PubMed:[10212285](#)). In esophageal epithelium, expressed in the basal, prickle and granular cell layers (PubMed:[22204712](#)). Up-regulated in a number of cancers cells and tissues.

Images



Western blot analysis of LOXL2 expression in MCF7 (A), K562 (B), Jurkat (C), mouse muscle (D) whole cell lysates. (Predicted band size: 86 kD; Observed band size: 53, 106 kD)

Immunohistochemical analysis of LOXL2 staining in human lung cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of LOXL2 staining in Hela cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a humidified chamber. Cells were washed with PBST and incubated with a AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark. Phalloidin - AF594 was used to stain Actin filaments (red). DAPI was used to stain the cell nuclei (blue).

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