

DDX5 (D15E10) XP[®] Rabbit mAb

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For Research Use Only. Not for Use in Diagnostic Procedures.

Applications:	Reactivity:	Sensitivity:	MW (kDa):	Source/Isotype:	UniProt ID:	Entrez-Gene Id:
W, IP, IF-IC, eCLIP	H M R Mk	Endogenous	68	Rabbit IgG	#P17844	1655

Product Usage Information**Application**

Western Blotting
Immunoprecipitation
Immunofluorescence (Immunocytochemistry)
eCLIP

Dilution

1:1000
1:50
1:100
1:200

For more information about the RBP-eCLIP service please visit Eclipsebio.

Storage

Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at -20°C. Do not aliquot the antibody.

For a carrier free (BSA and azide free) version of this product see product #75466.

Specificity/Sensitivity

DDX5 (D15E10) XP[®] Rabbit mAb recognizes endogenous levels of total DDX5 protein.

Species predicted to react based on 100% sequence homology

Bovine, Horse

Source / Purification

Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Ser498 of human DDX5 protein.

Background

DDX5 (DEAD box polypeptide 5), also known as p68, was first identified as a 68 kDa nuclear protein with similarity to translation initiation factor eIF-4A (1). DDX5 is a member of the DEAD box family of putative RNA helicases, defined by the presence of a conserved DEAD (Asp-Glu-Ala-Asp) motif that appears to function primarily in the regulation of RNA secondary structure. DDX5 exhibits ATP-dependent RNA helicase activity (2) and has been identified as a critical subunit of the DROSHA complex that regulates miRNA and rRNA processing (3,4). DDX5 may also regulate mRNA splicing (5) and has been shown to interact with HDAC1, where it can regulate promoter-specific transcription (6). DDX5 interacts with a diverse group of proteins, including Runx2, p53, Smad3, CBP, and p300 (7-10), suggesting an important role for DDX5 in a multitude of developmental processes. Notably, DDX5 may be involved in growth factor-induced epithelial mesenchymal transition (EMT). Phosphorylation of DDX5 at Tyr593 following PDGF stimulation was shown to displace Axin from β-catenin; this prevented phosphorylation of β-catenin by GSK-3β, leading to Wnt-independent nuclear translocation of β-catenin (11) and increased transcription of c-Myc, cyclin D1, and Snai1 (12,13).

Background References

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2. Hirling, H. et al. (1989) *Nature* 339, 562-4.
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4. Davis, B.N. et al. (2008) *Nature* 454, 56-61.
5. Camats, M. et al. (2008) *PLoS ONE* 3, e2926.
6. Wilson, B.J. et al. (2004) *BMC Mol Biol* 5, 11.
7. Jensen, E.D. et al. (2008) *J Cell Biochem* 103, 1438-51.
8. Bates, G.J. et al. (2005) *EMBO J* 24, 543-53.
9. Warner, D.R. et al. (2004) *Biochem Biophys Res Commun* 324, 70-6.
10. Rossow, K.L. and Janknecht, R. (2003) *Oncogene* 22, 151-6.
11. Yang, L. et al. (2006) *Cell* 127, 139-55.
12. Yang, L. et al. (2007) *J Biol Chem* 282, 16811-9.
13. Carter, C.L. et al. (2010) *Oncogene* 29, 5427-36.

Species Reactivity

Species reactivity is determined by testing in at least one approved application (e.g., western blot).

Western Blot Buffer

IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X TBS, 0.1% Tween@ 20 at 4°C with gentle shaking, overnight.

Applications Key

W: Western Blotting **IP:** Immunoprecipitation **IF-IC:** Immunofluorescence (Immunocytochemistry)
eCLIP: eCLIP

Cross-Reactivity Key

H: Human **M:** Mouse **R:** Rat **Mk:** Monkey

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