

YTHDC1 (E2P2I) Rabbit mAb



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Applications: W, IP	Reactivity: H M R Mk	Sensitivity: Endogenous	MW (kDa): 110	Source/Isotype: Rabbit IgG	UniProt ID: #Q96MU7	Entrez-Gene Id: 91746
Product Usage Information	2	Application Western Blotting Immunoprecipitation			Dilution 1:1000 1:50	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μ g/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		YTHDC1 (E2P2I) Rabbit mAb recognizes endogenous levels of total YTHDC1 protein.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Gly11 of human YTHDC1 protein.				
Background		YTH domain-containing protein 1 (YTHDC1) and YTH domain-containing protein 2 (YTHDC2) both belong to a family of proteins that bind to RNA. YTHDC1 and YTHDC2 both recognize and bind to N6-methyladenosine(m6A)-containing RNAs; binding is mediated through the YTH domains (1-3). m6A is a modification that is present at internal sites of mRNAs and some non-coding RNAs and plays a role in regulating mRNA splicing, processing, and stability. YTHDC1, also known as splicing factor YT521, regulates alternative splicing by functioning as a key regulator of exon-inclusion or exon-skipping. YTHDC1 promotes exon-inclusion by recruiting pre-mRNA splicing factor SRSF3 to regions containing m6A, while repressing exon-skipping by blocking SRSF10 binding to these same regions (2). Increased expression of YTHDC1 promotes malignant endometrial carcinoma (EC) through alternative splicing of vascular endothelial growth factor A (VEGF-A), resulting in an increase in VEGF-165 isoform and increased EC cell invasion (4). YTHDC2 functions to enhance the translation efficiency of target mRNAs and may play a role in spermatogenesis (5).				
Background R	eferences	1. Xu, C. et al. (2015) <i>J Biol Chem</i> 290, 24902-13. 2. Xiao, W. et al. (2016) <i>Mol Cell</i> 61, 507-19. 3. Xu, C. et al. (2014) <i>Nat Chem Biol</i> 10, 927-9. 4. Zhang, B. et al. (2015) <i>Tumor Biol</i> 37, 15543-9. 5. Hsu, P.J. et al. (2017) <i>Cell Res</i> 27, 1115-27.				
Consider Description		Consideration of the standard			d 1 1 1 1	

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

Cross-Reactivity Key H: Human M: Mouse R: Rat Mk: Monkey

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