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# Phospho-MOB1 (Thr35) (D2F10) Rabbit mAb



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

<b>Applications:</b> WB, IHC-P	Reactivity: H M R Mk	<b>Sensitivity:</b> Endogenous	MW (kDa): 24	Source/Isotype: Rabbit IgG	UniProt ID: #Q9H8S9, #Q7L9L4	<b>Entrez-Gene Id</b> 55233, 92597	
Product Usage Information	Aŗ	Application				Dilution	
	We	estern Blotting		1:1000			
	Im	munohistochemistry	(Paraffin)	1:50			
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 $\mu$ g/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at $-20^{\circ}$ C. Do not aliquot the antibody.					
Specificity / Sensitivity		Phospho-MOB1 (Thr35) (D2F10) Rabbit mAb recognizes endogenous levels of MOB1 protein only when phosphorylated at Thr35.					
Species predictereact based on 10	u	Hamster, Chicken, Xenopus, Zebrafish, Bovine, Horse, Guinea Pig					

### Source / Purification

sequence homology:

Monoclonal antibody is produced by immunizing animals with a synthetic phosphopeptide corresponding to residues surrounding Thr35 of human MOB1 protein.

## **Background**

MOB1 was first identified in yeast as a protein that binds to Mps with essential roles in the completion of mitosis and the maintenance of ploidy (1). Its *Drosophila* and mammalian homologs, Mats and MOB1, respectively, are involved in the Hippo signaling tumor suppressor pathway, which plays a critical role in organ size regulation and which has been implicated in cancer development (2-5). There are two MOB1 proteins in humans, MOB1A and MOB1B, that are encoded by two different genes but which have greater than 95% amino acid sequence identity (6). Both forms bind to members of the nuclear Dbf2-related (NDR) kinases, such as LATS1/2 and NDR1/2, thereby stimulating kinase activity (7-9). This binding is promoted by the phosphorylation of MOB1 at several threonine residues (e.g., Thr12, Thr35) by MST1 and/or MST2 (5,10).

Phosphorylation at Thr35 by MST1/2 stabilizes MOB1, enhancing its binding and regulation of LATS1 (5). The resultant increase in LATS1 kinase activity promotes inhibitory phosphorylation of the transcriptional co-activators YAP and TAZ (11,12), leading to changes in the expression of genes involved in cell cycle progression (13).

## **Background References**

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- 3. Saucedo, L.J. and Edgar, B.A. (2007) Nat Rev Mol Cell Biol 8, 613-21.
- 4. Harvey, K. and Tapon, N. (2007) Nat Rev Cancer 7, 182-91.
- 5. Zeng, Q. and Hong, W. (2008) Cancer Cell 13, 188-92.
- 6. Praskova, M. et al. (2008) Curr Biol 18, 311-21.
- 7. Devroe, E. et al. (2004) J Biol Chem 279, 24444-51.
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- 11. Zhao, B. et al. (2007) Genes Dev 21, 2747-61.
- 12. Lei, Q.Y. et al. (2008) Mol Cell Biol 28, 2426-36.
- 13. Hao, Y. et al. (2008) J Biol Chem 283, 5496-509.

## **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** 

WB: Western Blotting IHC-P: Immunohistochemistry (Paraffin)

**Cross-Reactivity Key** 

H: human M: mouse R: rat Hm: hamster Mk: monkey Vir: virus Mi: mink C: chicken Dm: D. melanogaster X: Xenopus Z: zebrafish B: bovine Dg: dog Pg: pig Sc: S. cerevisiae Ce: C. elegans Hr: horse

GP: Guinea Pig Rab: rabbit All: all species expected

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