

Phospho-TFEB (Ser122) Antibody



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Applications: W, IP	Reactivity: H	Sensitivity: Endogenous	MW (kDa): 70-80	Source/Isotype: Rabbit	UniProt ID: #P19484	Entrez-Gene Id: 7942
Product Usage Information	r	Application Western Blotting Immunoprecipitation			Dilution 1:1000 1:200	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 µg/ml BSA and 50% glycerol. Store at – 20°C. <i>Do not aliquot the antibody.</i>				
Specificity/Sensitivity		Phospho-TFEB (Ser122) Antibody recognizes endogenous levels of TFEB protein only when phosphorylated at Ser122.				
Species predicted to react based on 100% sequence homology		Mouse, Rat				
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a synthetic phosphopeptide corresponding to residues surrounding Ser122 of human TFEB protein. Antibodies are purified by peptide affinity chromatography.				
Background		Transcription factor EB (TFEB) is a member of the Myc-related, bHLH leucine-zipper family of transcription factors that drives the expression of a network of genes known as the Coordinated Lysosomal Expression and Regulation (CLEAR) network (1,2). TFEB specifically recognizes and binds regulatory sequences within the CLEAR box (GTCACGTGAC) of lysosomal and autophagy genes, resulting in the upregulated expression of genes involved in lysosome biogenesis and function, and regulation of autophagy (1,2). TFEB is activated in response to nutrient deprivation, stimulating translocation to the nucleus where it forms homo- or heterooligomers with other members of the microphthalmia transcription factor (MiTF) subfamily and resulting in upregulation of autophagosomes and lysosomes (3-5). Recently, it has been shown that TFEB is a component of mammalian target of rapamycin (mTOR) complex 1 (mTORC1), which regulates the phosphorylation and nuclear translocation of TFEB in response to cellular starvation and stress (6-9). During normal growth conditions, TFEB is phosphorylated at Ser211 in an mTORC1-dependent manner. Phosphorylation promotes association of TFEB with 14-3-3 family proteins and retention in the cytosol. Inhibition of mTORC1 results in a loss of TFEB phosphorylation, dissociation of the TFEB/14-3-3 complex, and rapid transport of TFEB to the nucleus where it increases transcription of CLEAR and autophagy genes (10). TFEB has also been shown to be activated in a nutrient-dependent manner by p42 MAP kinase (Erk2). TFEB is phosphorylated at Ser142 by Erk2 in response to nutrient deprivation, resulting in nuclear localization and activation, and indicating that pathways other than mTOR contribute to nutrient sensing via TFEB (3). Additional studies have also identified phosphorylation of TFEB at Ser122 that is dependent on mTORC1 (11).				
Background References		 Sardiello, M. et al. (2009) Science 325, 473-7. Sardiello, M. and Ballabio, A. (2009) Cell Cycle 8, 4021-2. Settembre, C. et al. (2011) Science 332, 1429-33. David, R. (2011) Nat Rev Mol Cell Biol 12, 404. Cuervo, A.M. (2011) Science 332, 1392-3. Peña-Llopis, S. et al. (2011) EMBO J 30, 3242-58. Settembre, C. and Ballabio, A. (2011) Autophagy 7, 1379-81. Peña-Llopis, S. and Brugarolas, J. (2011) Cell Cycle 10, 3987-8. Settembre, C. et al. (2012) EMBO J 31, 1095-108. Martina, J.A. et al. (2012) Autophagy 8, 903-14. 				

11. Vega-Rubin-de-Celis, S. et al. (2017) *Autophagy* 13, 464-72.

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

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