

## **BRF1/2 Antibody**



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

Applications: W	Reactivity: H M R Mk	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 40 to 50, 62	Source/Isotype: Rabbit	<b>UniProt ID:</b> #P47974, #Q07352	<b>Entrez-Gene Id</b> : 678, 677
Product Usage Information		<b>Application</b> Western Blotting			<b>Dilution</b> 1:1000	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 $\mu$ g/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		This antibody detects endogenous levels of total BRF1 and BRF2 proteins.				
Species predicted to react based on 100% sequence homology		Chicken, Bovine				
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to the carboxy terminus of human BRF1 protein. Antibodies are purified by protein A and peptide affinity chromatography.				
Background		ZFP36L1, also known as butyrate response factor-1 (BRF1), and ZFP36L2, also known as butyrate response factor-2 (BRF2), both belong to the TIS11 family of CCCH zinc-finger proteins (1). This family of proteins, which also includes tristetraprolin (TTP), bind to AU-rich elements (AREs) found in the 3'-untranslated regions of mRNAs and promote deadenylation and rapid degradation by the exosome (2,3). These proteins play a critical role in cell growth control by regulating the mRNA turnover of multiple cytokines, growth factors, and cell cycle regulators, including GM-CSF, TNFα, IL-2, IL-3, and IL-6 (4,5). Deregulated ARE-mRNA stability can contribute to both inflammation and oncogenic transformation (6-8). Insulin-induced stabilization of ARE-containing transcripts is mediated by Akt/PKB phosphorylation of ZFP36L1 at Ser92, which results in binding by 14-3-3 protein and inactivation of ZFP36L1 (9). ZFP36L1 and L2 have also been shown to promote cell quiescence in developing B lymphocytes, promoting VDJ recombination (10).				
Background References		1. Varnum, B.C. et al. (1991) <i>Mol. Cell. Biol.</i> 11, 1754-1758. 2. Stoecklin, G. et al. (2002) <i>EMBO J.</i> 21, 4709-4718. 3. Lykke-Andersen, J. and Wagner, E. (2005) <i>Genes Dev.</i> 19, 351-361. 4. Stoecklin, G. et al. (2000) <i>Mol. Cell. Biol.</i> 20, 3753-3763. 5. Stoecklin, G. et al. (2001) <i>RNA</i> 7, 1578-1588. 6. Schuler, G.D. and Cole, M.D. (1988) <i>Cell</i> 55, 1115-1122. 7. Nair, A.P. et al. (1994) <i>Nature</i> 369, 239-242. 8. Carballo, E. et al. (1998) <i>Science</i> 281, 1001-1005. 9. Schmidlin, M. et al. (2004) <i>EMBO J.</i> 23, 4760-4769. 10. Galloway, A. et al. (2016) <i>Science</i> 352, 453-9.				
Species Reactivi	ity	Species reactivity is d	etermined by testin	g in at least one appro	ved application (e.g., w	estern 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

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

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