HIF1 bata Monoclonal Antibody

catalog number: E-AB-22189



Note: Centrifuge before opening to ensure complete recovery of vial contents.

Description			
Reactivity	Mouse	Mouse	
Immunogen	Recombinant Protein	Recombinant Protein	
Host	Mouse	Mouse	
Isotype	IgG	IgG	
Clone	4C5	4C5	
Purification	Protein A purification		
Conjugation	Unconjugated	Unconjugated	
buffer	Phosphate buffered solu	Phosphate buffered solution, pH 7.4, containing 0.05% stabilizer, 0.5% protein	
	protectant and 50% glyc	protectant and 50% glycerol.	
Applications	Recommended Dilu	Recommended Dilution	
WB	1:1000-2000	1:1000-2000	
IHC	1:100-200		
Data			
14. Western Blot analysi Monoclonal Ar	s of Mouse brain using HIF1 bata tibody at dilution of 1:2000.	Immunohistochemistry of paraffin-embedded Mouse brain tissue using HIF1 bata Monoclonal Antibody at dilution of	
Observed-MV:87 kDa		1:200.	
Preparation & Storage			
Storage	Store at -20°C Valid for 1	Store at -20°C Valid for 12 months. Avoid freeze / thaw cycles.	
Shipping	The product is shipped v	The product is shipped with ice pack, upon receipt, store it immediately at the	
	temperature recommende	temperature recommended.	

Background

Hypoxia-inducible factor 1 (HIF1) is a heterodimeric transcription factor that plays a critical role in the cellular response to hypoxia (1). The HIF1 complex consists of two subunits, HIF-1 α and HIF-1 β , which are basic helix-loop-helix proteins of the PAS (Per, ARNT, Sim) family (2). HIF1 regulates the transcription of a broad range of genes that facilitate responses to the hypoxic environment, including genes regulating angiogenesis, erythropoiesis, cell cycle, metabolism and apoptosis. The widely expressed HIF-1 α is typically degraded rapidly in normoxic cells by the ubiquitin/proteasomal pathway. Under normoxic conditions, HIF-1 α is proline hydroxylated leading to a conformational change that promotes binding to the von Hippel Lindau protein (VLH) E3 ligase complex, ubiquitination and proteasomal degradation follows (3,4). Both hypoxic conditions and chemical hydroxylase inhibitors (such as desferrioxamine and cobalt) inhibit HIF-1 α degradation and lead to its stabilization. In addition, HIF-1 α can be induced in an oxygen-independent manner by various cytokines through the PI3K-AKT-mTOR pathway (5-7).HIF-1 β is also known as AhR nuclear translocator (ARNT) due to its ability to partner with the aryl hydrocarbon receptor (AhR) to form a heterodimeric transcription factor complex (8). Together with AhR, HIF-1 β plays an important role in xenobiotics metabolism (8).

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