

EPAS1 Antibody (N-term)
Affinity Purified Rabbit Polyclonal Antibody (Pab)
Catalog # AP10707a**Specification**

EPAS1 Antibody (N-term) - Product Information

Application	FC, WB,E
Primary Accession	O99814
Other Accession	NP_001421.2
Reactivity	Human
Host	Rabbit
Clonality	Polyclonal
Isotype	Rabbit IgG
Calculated MW	96459
Antigen Region	124-150

EPAS1 Antibody (N-term) - Additional Information**Gene ID** 2034**Other Names**

Endothelial PAS domain-containing protein 1, EPAS-1, Basic-helix-loop-helix-PAS protein MOP2, Class E basic helix-loop-helix protein 73, bHLHe73, HIF-1-alpha-like factor, HLF, Hypoxia-inducible factor 2-alpha, HIF-2-alpha, HIF2-alpha, Member of PAS protein 2, PAS domain-containing protein 2, EPAS1, BHLHE73, HIF2A, MOP2, PASD2

Target/Specificity

This EPAS1 antibody is generated from rabbits immunized with a KLH conjugated synthetic peptide between 124-150 amino acids from the N-terminal region of human EPAS1.

Dilution

FC~~1:10~50

WB~~1:1000

E~~Use at an assay dependent concentration.

Format

Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This antibody is purified through a protein A column, followed by peptide affinity purification.

Storage

Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.

Precautions

EPAS1 Antibody (N-term) is for research use only and not for use in diagnostic or therapeutic procedures.

EPAS1 Antibody (N-term) - Protein Information

Name EPAS1**Synonyms** BHLHE73, HIF2A, MOP2, PASD2

Function Transcription factor involved in the induction of oxygen regulated genes. Heterodimerizes with ARNT; heterodimer binds to core DNA sequence 5'-TACGTG-3' within the hypoxia response element (HRE) of target gene promoters (By similarity). Regulates the vascular endothelial growth factor (VEGF) expression and seems to be implicated in the development of blood vessels and the tubular system of lung. May also play a role in the formation of the endothelium that gives rise to the blood brain barrier. Potent activator of the Tie-2 tyrosine kinase expression. Activation requires recruitment of transcriptional coactivators such as CREBBP and probably EP300. Interaction with redox regulatory protein APEX1 seems to activate CTAD (By similarity).

Cellular Location

Nucleus {ECO:0000250|UniProtKB:P97481, ECO:0000255|PROSITE-ProRule:PRU00981}. Nucleus speckle {ECO:0000250|UniProtKB:P97481}. Note=Colocalizes with HIF3A in the nucleus and speckles. {ECO:0000250|UniProtKB:P97481}

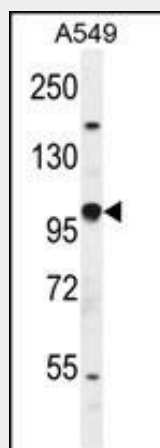
Tissue Location

Expressed in most tissues, with highest levels in placenta, lung and heart. Selectively expressed in endothelial cells

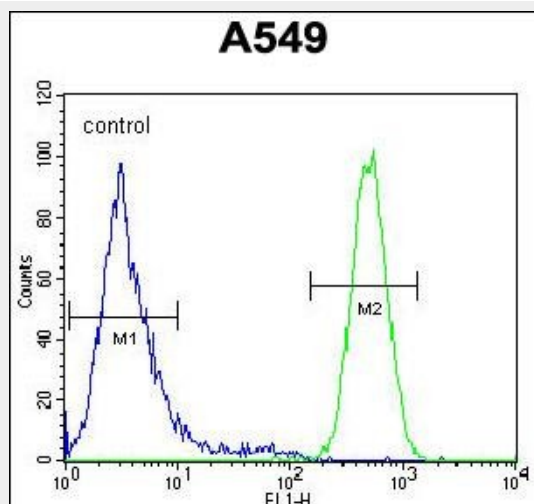
EPAS1 Antibody (N-term) - Protocols

Provided below are standard protocols that you may find useful for product applications.

- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

EPAS1 Antibody (N-term) - Images

EPAS1 Antibody (N-term) (Cat. #AP10707a) western blot analysis in A549 cell line lysates (35ug/lane). This demonstrates the EPAS1 antibody detected the EPAS1 protein (arrow).



EPAS1 Antibody (N-term) (Cat. #AP10707a) flow cytometric analysis of A549 cells (right histogram) compared to a negative control cell (left histogram). FITC-conjugated goat-anti-rabbit secondary antibodies were used for the analysis.

EPAS1 Antibody (N-term) - Background

This gene encodes a transcription factor involved in the induction of genes regulated by oxygen, which is induced as oxygen levels fall. The encoded protein contains a basic-helix-loop-helix domain protein dimerization domain as well as a domain found in proteins in signal transduction pathways which respond to oxygen levels. Mutations in this gene are associated with erythrocytosis familial type 4.

EPAS1 Antibody (N-term) - References

Bailey, S.D., et al. Diabetes Care 33(10):2250-2253(2010)
Yi, X., et al. Science 329(5987):75-78(2010)
Bougatef, F., et al. PLoS ONE 5 (8), E12265 (2010) :
Hosseini Ghaderian, S.M., et al. Pathology 42(5):446-453(2010)
Mowat, F.M., et al. PLoS ONE 5 (6), E11103 (2010) :