

## Anti-Human HIF-2-ALPHA [Polyclonal]

Catalog No.	Description		
ARB28-5R	6 ml of Ready-to-Use Antibody for use with BioGenex Super Sensitive <sup>TM</sup> Detection Systems OR equivalent detection system		
ARB28-10R  10 ml of Ready-to-Use Antibody in a barcode labeled vial for use with BioGen Super Sensitive TM Detection Systems and i6000 TM Automated Staining Systems			
PUB28-UP	1 ml of Concentrated Antibody for use with BioGenex Super Sensitive <sup>TM</sup> Detection Systems OR equivalent detection system		
PUB28-5UP  0.5 ml of Concentrated Antibody for use with BioGenex Super Sensitive TM Detect Systems OR equivalent detection system			
AWB28-YCD	Ready-to-Use Antibody in Barcode labeled vial for use on the Xmatrx <sup>®</sup> Elite/Ultra Staining System, 160 tests		
AWB28-50D Ready-to-Use Antibody in Barcode lad vial for use on the Xmatrx® Elite/Ultra Staining System, 50 tests			

Clone	Species	Ig Class
Polyclonal	Rabbit	IgG

#### **Intended Use**

For In Vitro Diagnostic Use. This antibody is designed for the specific localization of HIF-2-alpha in formalin-fixed, paraffinembedded (FFPE) tissue sections. Evaluation must be performed by a qualified pathologist.

### **Summary and Explanation**

HIF-2-alpha (also known as EPAS1) is a transcription factor involved in the induction of genes regulated by oxygen. It shares 48% sequence identity with HIF1-alpha (HIF1A). Like HIF1A, HIF-2-alpha regulates gene expression in response to hypoxia. It also regulates the vascular endothelial growth factor (VEGF) expression and seems to be implicated in the development of blood vessels and the tubular system of the lungs. HIF2A is expressed at relatively higher levels in villus sections of placenta and in lung samples.

#### **Storage and Handling**

Store at 2-8°C. Fresh dilutions, if required, should be prepared prior to use and are stable and steady for up to one day at room temperature (20-26°C). Diluted antibody preparations can be refrigerated or frozen for extended shelf life.

# Emergo Europe, Prinsessegracht 20, 2514AP The Hague, The Netherlands

#### **Principles of the Procedure**

Antigen detection by immunohistochemistry (IHC) is a two-step process wherein the primary antibody binds to the antigen of interest and that binding is detected by a chromogen. The primary antibody may be used in IHC using manual techniques or BioGenex Automated Staining System. Positive and negative controls should always be run simultaneously with all patient specimens.

#### **Reagents Provided**

Rabbit Polyclonal Antibody to HIF-2-alpha is affinity purified and diluted in PBS, pH 7.2, containing 1% BSA and 0.09% sodium azide.

#### **Dilution of Primary Antibody**

BioGenex Ready-to-Use antibodies have been optimized for use with the recommended BioGenex Detection System and should not require further dilution.

BioGenex concentrated antibodies must be diluted in accordance with the recommended protocol when used with the recommended BioGenex Detection System.

#### **Recommended Protocol**

Refer to the following table for conditions specifically recommended for this antibody. Refer to the BioGenex website for guidance on specific staining protocols or other requirements.

Parameter	BioGenex Recommendations	
Control Tissue	Lung Tissues as available with Biogenex FB-B28P* & FG-B28P*	
Recommended Dilution for Concentrated Antibody	1:20-50 in HK941	
Recommended Pretreatment (Manual/i6000)**	EZ-AR2 (HK522-XAK)	
Recommended Pretreatment (Xmatrx)	EZ-AR2 Elegance (HX032-YCD)	
Antibody Incubation (Manual/i6000)	30-60 min at RT	
Antibody Incubation (Xmatrx)	30-60 min at 25°C	
Detection System for Manual, Xmatrx & i6000 systems***	Use BioGenex Two-Step <b>OR</b> One-Step Super Sensitive <sup>™</sup> Polymer-HRP IHC Detection System/DAB; see p. 2 for more information	

\*FB: positive control barrier slides, FG: positive control nonbarrier slides. Xmatrx requires barrier slides.

\*\*Pretreatment times will vary based on individual microwave power. \*\*\*For automation systems (Xmatrx-Elite, Xmatrx-Ultra & i6000 Diagnostics), refer to the factory protocols provided with the instrument.

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CE	Emergo Europe, Prinsessegracht 20, 2514AP The Hague, The Netherlands
EC REF	

Detection	Two-Step	One-Step	Link and
System	HRP Kit	HRP Kit	Label Kit
Manual	QD440-XAKE (1000 Test) QD430-XAKE (1000 Test)	QD630-XAKE (1000 Test)	QP300-XAKE (1000 Test)
Manuai	QD420-YIKE (500 Test) QD400-60KE (60 Test)	QD620-XAKE (500 Test)	QP900-9LE (500 Test)
Xmatrx -	QD550-YCDE	QD610-YADE	N/A
Automation	(200 Test)	(200 Test)	
i6000 -	QD410-YAXE	QD610-YAXE	N/A
Automation	(200 Test)	(200 Test)	
For more information, visit www.biogenex.com.			

#### **Precautions**

This product contains sodium azide at concentrations of less than 0.1%. Sodium azide is not classified as a hazardous chemical at the product concentrations, but proper handling protocols should be observed. For more information, a Safety Data Sheet (SDS) for sodium azide is available upon request. Dispose of unused reagents according to Local, State and Federal Regulations. Wear suitable Personal Protective Equipment, do not pipette reagents by mouth, and avoid contact of reagents and specimens with skin and mucous membranes. If reagents or specimens come in contact with sensitive area, wash with copious amounts of water.

### **Quality Control**

Refer to BioGenex detection system documents for guidance on general quality control procedures.

#### **Troubleshooting**

Refer to the troubleshooting section in the documentation for BioGenex Detection Systems (or equivalent detection systems) for remedial actions on detection system related issues, or contact BioGenex Technical Support Department at 1-800-421-4149 or <a href="mailto:support@biogenex.com">support@biogenex.com</a> or your local distributor to report unusual staining.

#### **Expected Results**

This antibody stains membrane/cytoplasm in positive cells in formalin-fixed, paraffin embedded tissue sections. An example image of a tissue section stained with this antibody can be found on the product page on the BioGenex website. Interpretation of the staining result is solely the responsibility of the user. Experimental results should be confirmed by a medically-established diagnostic product or procedure.

#### **Limitations of the Procedure**

Improper tissue handling and processing prior to immunostaining can lead to inconsistent results. Variations in embedding and fixation or the nature of the tissue may lead to variations in results. Endogenous peroxidase activity or pseudo peroxidase activity in erythrocytes and tissue biotin may result in non-specific staining based on the detection system employed. Tissues containing Hepatitis B Surface Antigen (HBsAg) may give false positive with horseradish peroxidase systems. Improper counterstaining and mounting may compromise the interpretation of results.

#### **Bibliography**

- Highet AR, Khoda SM, Buckberry S et al. Hypoxia induced HIF-1/HIF-2 activity alters trophoblast transcriptional regulation and promotes invasion. Eur. J. Cell Biol. 2015 Dec 01
- 2. Chen C, Ma Q, Ma X et al. Association of Elevated HIF-2a Levels with Low Beclin 1 Expression and Poor Prognosis in Patients with Chondrosarcoma. Ann Surg Oncol. 2011 Feb 16.
- Groenman F, Rutter M, Caniggia I et al. Hypoxia-inducible factors in the first trimester human lung. J Histochem Cytochem 2007.
- Nakazawa, MS. Et al. Epigenetic re-expression of HIF-2α suppresses soft tissue sarcoma growth. Nat Commun. 2016 Feb 3; 7:10539.

2°C 8°C	Temperature Limitation	IVD	In Vitro Diagnostic Medical Device
$\boxtimes$	Use By Date	LOT	Batch Code
NON	Non-Sterile	i	Consult Instructions for Use
EC REP	Representative in the European Community		Manufacturer

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