Anti-HIF1 alpha Antibody

ER1802-41



| Product Type: Species reactivity: | Rabbit polyclonal IgG, primary antibodies Human, Mouse, Rat |
|---|---|
| Applications: | WB, IF-Cell, FC |
| Molecular Wt: | Predicted band size: 92 kDa |
| Description: | Cell growth and viability is compromised by oxygen deprivation (hypoxia). Hypoxia- inducible factors, including HIF-1 α , HIF-1 β (also designated Arnt 1), EPAS-1 (also designated HIF-2 α) and HIF-3 α , induce glycolysis, erythropoiesis and angiogenesis in order to restore oxygen homeostasis. Hypoxia-inducible factors are members of the Per- Arnt-Sim (PAS) domain transcription factor family. In response to hypoxia, HIF-1 α is upregulated and forms a heterodimer with Arnt 1 to form the HIF-1 complex. The HIF-1 complex recognizes and binds to the hypoxia responsive element (HRE) of hypoxia- inducible genes, thereby activating transcription. Hypoxia-inducible expression of some genes such as Glut-1, p53, p21 or Bcl-2, is HIF-1 α independent, whereas expression of others, such as p27, GADD 153 or HO-1, is HIF-1 α independent. EPAS-1 and HIF-3 α have also been shown to form heterodimeric complexes with Arnt 1 in response to hypoxia. |
| lmmunogen: | Synthetic peptide within C-terminal human HIF1 alpha. |
| Subcellular location: | Cytoplasm. Nucleus. |
| Database links: | SwissProt: Q16665 Human Q61221 Mouse |
| Recommended Dilutions: WB IF-cell FC | 1:500-2000 1:100 1µg/Test: 1:5000-10000 |
| Storage Buffer: | 1*PBS (pH7.4), 0.2% BSA, 50% Glycerol. Preservative: 0.05% Sodium Azide. |
| Storage Instruction: | Shipped at 4 $^\circ\!\!\mathbb{C}$. Store at -20 $^\circ\!\!\mathbb{C}$ for one year. Avoid repeated freeze/thaw cycles. |
| Purity: | Protein A affinity purified. |

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Applications:WB=Western blot IHC-P=Immunohistochemistry (paraffin) IF-Cell=Immunofluorescence (Cell) IF-Tissue=Immunofluorescence (Tissue) FC=Flow cytometry IP=Immunoprecipitation

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Fig1: 50 ng RHHIF-1 Alpha-Trx-His protein per lane probed with HIF-1 Alpha polyclonal antibody respectively, unconjugated (ER1802-41) at 1:1000 dilution and 4 $^{\circ}$ C overnight incubation. Followed by corresponding conjugated secondary antibody incubation at r.t. for 60 min.



Fig2: 25 ug total protein per lane of various lysates (see on figure) probed with HIF-1 Alpha polyclonal antibody, unconjugated (ER1802-41) at 1:1000 dilution and 4°C overnight incubation. Followed by conjugated secondary antibody incubation at r.t. for 60 min.



Fig3: Hela cell; 4% Paraformaldehyde-fixed; Triton X-100 at room temperature for 20 min; Blocking buffer (normal goat serum, C-0005) at 37°C for 20 min; Antibody incubation with (HIF-1 Alpha) polyclonal Antibody, Unconjugated (ER1802-41) 1:100, 90 minutes at 37°C; followed by a conjugated Goat Anti-Rabbit IgG antibody at 37°C for 90 minutes, DAPI (blue, C02-04002) was used to stain the cell nuclei.



Fig4: Hela cell; 4% Paraformaldehyde-fixed; Triton X-100 at room temperature for 20 min; Blocking buffer (normal goat serum, C-0005) at 37°C for 20 min; Antibody incubation with (HIF-1 Alpha) polyclonal Antibody, Unconjugated (ER1802-41) 1:100, 90 minutes at 37°C; followed by a conjugated Goat Anti-Rabbit IgG antibody at 37°C for 90 minutes, DAPI (blue, C02-04002) was used to stain the cell nuclei.

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Fig5: Blank control (blue line): Hela (blue). Primary Antibody (green line): Rabbit Anti- HIF-1 Alpha antibody (ER1802-41) Dilution: 1µg /10⁶ cells; Isotype Control Antibody (orange line): Rabbit IgG . Secondary Antibody (white blue line): Goat anti-rabbit IgG-FITC Dilution: 1µg /test. Protocol The cells were fixed with 80% methanol (5 min at -20 $^{\circ}$ C) and then permeabilized with 0.1% PBS-Tween for 20 min at room temperature. Cells stained with Primary Antibody for 30 min at room temperature. The cells were then incubated in 1 X PBS/2%BSA/10% goat serum to block non-specific protein-protein interactions followed by the antibody for 15 min at room temperature. The secondary antibody used for 40 min at room temperature. Acquisition of 20,000 events was performed.

Note: All products are "FOR RESEARCH USE ONLY AND ARE NOT INTENDED FOR DIAGNOSTIC OR THERAPEUTIC USE".

Background References

- 1. Bhattacharya S et al. Functional role of p35srj, a novel p300/CBP binding protein, during transactivation by HIF-1. Genes Dev 13:64-75 (1999).
- 2. Masson N et al. Independent function of two destruction domains in hypoxia-inducible factor-alpha chains activated by prolyl hydroxylation. EMBO J 20:5197-5206 (2001).

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