

Product Datasheet

BICD1 Antibody - BSA Free

NBP1-78735

Unit Size: 100 ul

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Updated 2/21/2025 v.20.1

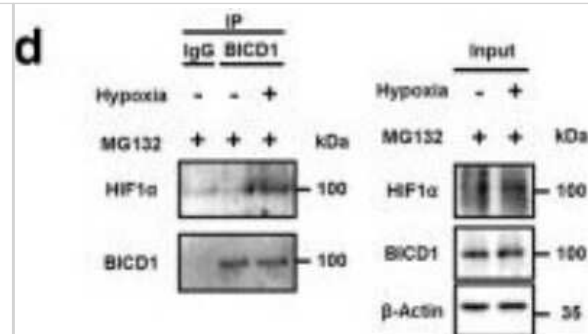
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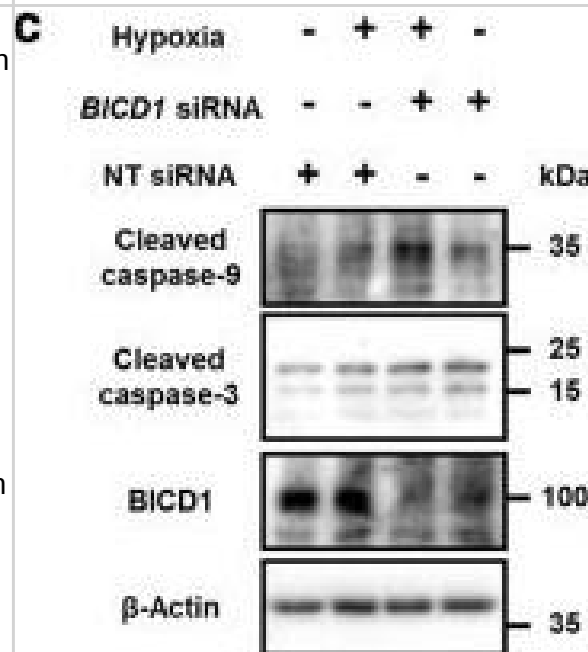
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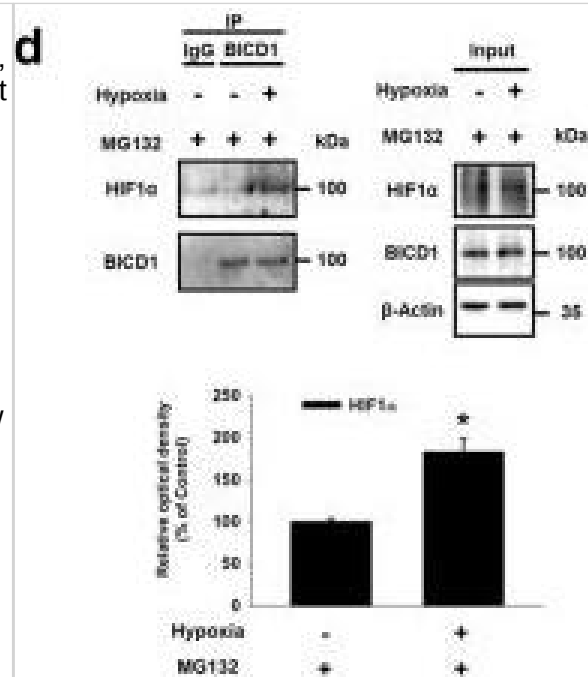
Western Blot: BICD1 Antibody [NBP1-78735] - Effect of hypoxia on the interaction between HIF-1 alpha and BICD1. Cells were pretreated with MG132 (1 μ M) for 30 min prior to hypoxia treatment for 24 h. Co-immunoprecipitation of HIF-1 alpha with IgG and BICD1 were shown in left panel. Total protein expressions in lysate were shown in right panel. $n = 3$. Image collected and cropped by CiteAb from the following publication (<https://www.nature.com/articles/s41418-018-0241-1>), licensed under a CC-BY license.



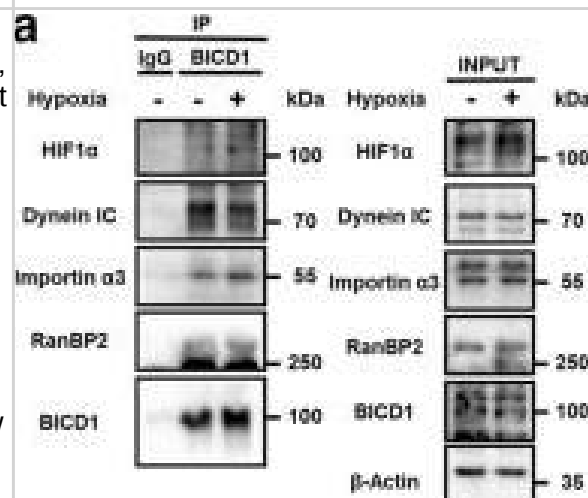
Western Blot: BICD1 Antibody [NBP1-78735] - Role of BICD1 in UCB-MSCs survival under hypoxia. a, b The UCB-MSCs were transfected with BICD1 or NT siRNA for 24 h prior to hypoxia treatment. a Cells were exposed to various durations of hypoxia (0–72 h). Cell viability of UCB-MSCs was measured by WST-1 cell viability assay. $n = 8$. b Cells were incubated in normoxia or hypoxia conditions for 72 h. Representative images of experimental groups at 0, 24, 48, & 72 h of normoxia or hypoxia incubation are presented. Red-marked cells in representative images indicate PI-positive cells. $n = 4$. Scale bars are 50 μ m (Magnification, $\times 200$). c BICD1 or NT siRNA-transfected cells were incubated in hypoxia for 24 h. Cleaved caspase-9, cleaved caspase-3, & β -Actin protein expressions were detected by western blot analysis. $n = 4$. d Cells were transfected with BICD1 or NT siRNA for 24 h prior to hypoxia treatment for 48 h. The percentages of apoptotic cells were analyzed by Annexin V/PI analysis, measured by flowcytometer. Annexin V-positive cells were considered as apoptotic cells. $n = 4$. Quantitative data are presented as a mean $\pm$ S.E.M. All blot images are representative. * $p < 0.05$ vs. normoxia control with NT siRNA transfection, # $p < 0.05$ vs. hypoxia with NT siRNA transfection. e The UCB-MSCs were transfected with pcDNA3.1/BICD1-cEGFP vector, pcDNA3.1/cEGFP vector, HIF1A siRNA or NT siRNA for 24 h prior to hypoxia treatment for 72 h. Cell viability data are presented as a mean $\pm$ S.E.M. $n = 5$. * $p < 0.05$ vs. normoxia control with pcDNA3.1/cEGFP vector & NT siRNA cotransfection, # $p < 0.05$ vs. hypoxia control with pcDNA3.1/cEGFP vector & NT siRNA cotransfection, @ $p < 0.05$ vs. hypoxia control with pcDNA3.1/BICD1-cEGFP vector & NT siRNA cotransfection. Image collected & cropped by CiteAb from the following publication (<https://pubmed.ncbi.nlm.nih.gov/30464225>), licensed under a CC-BY license. Not internally tested by Novus Biologicals.



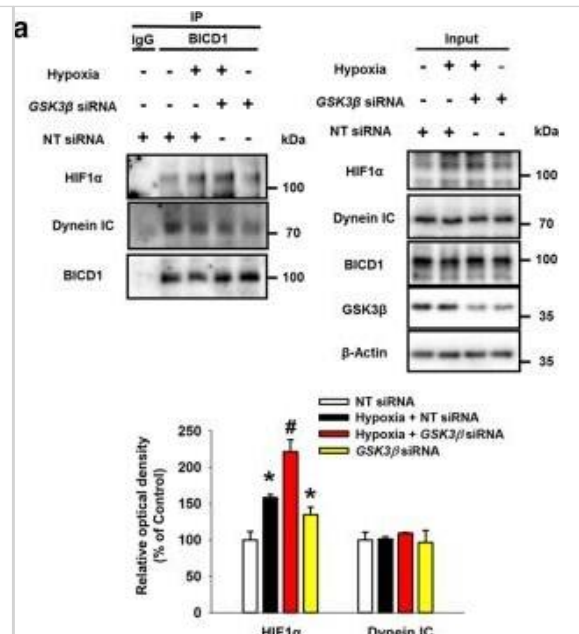
Western Blot: BICD1 Antibody [NBP1-78735] - Effect of hypoxia on the interaction between HIF1 α & BICD1. a Co-immunoprecipitation of HIF1 α , Dynein IC, Importin α 3, RanBP2 w/ IgG & BICD1 antibodies shown in left panel. Total protein expressions in lysate shown in right panel. n = 3. b Cells immunostained w/ HIF1 α & BICD1-specific antibodies. Scale bars are 8 μ m (Magnification, $\times$ 1,000). White arrow heads indicate co-localization of HIF1 α w/ BICD1. c Interaction between HIF1 α & BICD1 (HIF1 α /BICD1, red) analyzed by PLA. Scale bars are 8 μ m (Magnification, $\times$ 1,000). n = 6. *p < 0.05 vs. normoxia control. d-f Cells pretreated w/ MG132 (1 μ M) for 30 min prior to hypoxia treatment for 24 h. d Co-immunoprecipitation of HIF1 α w/ IgG & BICD1 shown in left panel. Total protein expressions in lysate shown in right panel. n = 3. e Cells immunostained w/ HIF1 α & BICD1-specific antibodies. White arrow heads indicate co-localization of HIF1 α w/ BICD1 in MG132-pretreated UCB-MSCs. Scale bars are 8 μ m (Magnification, $\times$ 1,000). n = 5. f HIF1 activity measured by dual luciferase reporter assay. n = 6. *p < 0.05 vs. normoxia control w/ MG132 pretreatment. g, h Cells transfected w/ BICD1 or NT siRNA for 24 h prior to hypoxia treatment for 24 h. g Co-immunoprecipitation of Dynein IC w/ IgG & HIF1 α antibodies shown in left panel. Total protein expressions in lysate shown in right panel. n = 3. h Interaction between HIF1 α & Dynein IC (HIF1 α /Dynein IC, red) analyzed by PLA. Scale bars are 8 μ m (Magnification, $\times$ 1,000). Quantitative data are presented as a mean $\pm$ S.E.M. All blots & immunofluorescence images are representative. *p < 0.05 vs. normoxia control w/ NT siRNA transfection, #p < 0.05 vs. hypoxia w/ NT siRNA transfection Image collected & cropped by CiteAb from the following publication (<https://pubmed.ncbi.nlm.nih.gov/30464225>), licensed under a CC-BY license. Not internally tested by Novus Biologicals.



Western Blot: BICD1 Antibody [NBP1-78735] - Effect of hypoxia on the interaction between HIF1 α & BICD1. a Co-immunoprecipitation of HIF1 α , Dynein IC, Importin α 3, RanBP2 w/ IgG & BICD1 antibodies shown in left panel. Total protein expressions in lysate shown in right panel. n = 3. b Cells immunostained w/ HIF1 α & BICD1-specific antibodies. Scale bars are 8 μ m (Magnification, $\times$ 1,000). White arrow heads indicate co-localization of HIF1 α w/ BICD1. c Interaction between HIF1 α & BICD1 (HIF1 α /BICD1, red) analyzed by PLA. Scale bars are 8 μ m (Magnification, $\times$ 1,000). n = 6. *p < 0.05 vs. normoxia control. d-f Cells pretreated w/ MG132 (1 μ M) for 30 min prior to hypoxia treatment for 24 h. d Co-immunoprecipitation of HIF1 α w/ IgG & BICD1 shown in left panel. Total protein expressions in lysate shown in right panel. n = 3. e Cells immunostained w/ HIF1 α & BICD1-specific antibodies. White arrow heads indicate co-localization of HIF1 α w/ BICD1 in MG132-pretreated UCB-MSCs. Scale bars are 8 μ m (Magnification, $\times$ 1,000). n = 5. f HIF1 activity measured by dual luciferase reporter assay. n = 6. *p < 0.05 vs. normoxia control w/ MG132 pretreatment. g, h Cells transfected w/ BICD1 or NT siRNA for 24 h prior to hypoxia treatment for 24 h. g Co-immunoprecipitation of Dynein IC w/ IgG & HIF1 α antibodies shown in left panel. Total protein expressions in lysate shown in right panel. n = 3. h Interaction between HIF1 α & Dynein IC (HIF1 α /Dynein IC, red) analyzed by PLA. Scale bars are 8 μ m (Magnification, $\times$ 1,000). Quantitative data are presented as a mean $\pm$ S.E.M. All blots & immunofluorescence images are representative. *p < 0.05 vs. normoxia control w/ NT siRNA transfection, #p < 0.05 vs. hypoxia w/ NT siRNA transfection Image collected & cropped by CiteAb from the following publication (<https://pubmed.ncbi.nlm.nih.gov/30464225>), licensed under a CC-BY license. Not internally tested by Novus Biologicals.



Western Blot: BICD1 Antibody [NBP1-78735] - Involvement of Akt/GSK3 β pathway in BICD1-mediated HIF1 α nuclear translocation.
a–e The UCB-MSCs were transfected with GSK3 β or NT siRNA for 24 h prior to hypoxia treatment for 24 h. a Co-immunoprecipitation of HIF1 α & Dynein IC with IgG & BICD1 antibodies were shown in left panel. Total protein expressions in lysate were shown in right panel. n = 3. b Interaction between HIF1 α & BICD1 (HIF1 α /BICD1, red) was analyzed by PLA assay. Scale bars are 8 μ m (Magnification, $\times$ 1,000). n = 5. c HIF1 α , Lamin A/C, & α -Tubulin in cytosolic & nuclear fractionized samples were detected by western blot. n = 4. d Cells were immunostained with HIF1 α -specific antibody. Scale bars are 8 μ m (Magnification, $\times$ 1,000). n = 4. e HIF1 activities in NT or GSK3 β siRNA-transfected cells were analyzed by dual luciferase reporter assay. n = 6. Quantitative data are presented as a mean $\pm$ S.E.M. All blots & immunofluorescence images are representative. *p < 0.05 vs. normoxia control with NT siRNA transfection, #p < 0.05 vs. hypoxia with NT siRNA transfection. Image collected & cropped by CiteAb from the following publication (<https://pubmed.ncbi.nlm.nih.gov/30464225>), licensed under a CC-BY license. Not internally tested by Novus Biologicals.



Publications

Jiang Y, Yao B, Chen T et al. BICD1 functions as a prognostic biomarker and promotes hepatocellular carcinoma progression *Pathol. Res. Pract.* 2020-02-12 [PMID: 32088084] (WB, Human)

Lee HJ, Jung YH, Choi GE et al. O-cyclic phytosphingosine-1-phosphate stimulates HIF1 alpha-dependent glycolytic reprogramming to enhance the therapeutic potential of mesenchymal stem cells *Cell Death Dis* 2019-08-05 [PMID: 31383843] (WB)

Lee HJ, Jung YH, Oh JY et al. BICD1 mediates HIF1a nuclear translocation in mesenchymal stem cells during hypoxia adaptation. *Cell Death Differ.* 2018-11-21 [PMID: 30464225] (WB, Human)



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