

Product Datasheet

Erythropoietin R Antibody - BSA Free NBP1-19388

Unit Size: 0.1 ml

Store at 4C short term. Aliquot and store at -20C long term. Avoid freeze-thaw cycles.

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NBP1-19388

Erythropoietin R Antibody - BSA Free

Product Information	
Unit Size	0.1 ml
Concentration	1.0 mg/ml
Storage	Store at 4C short term. Aliquot and store at -20C long term. Avoid freeze-thaw cycles.
Clonality	Polyclonal
Preservative	0.02% Sodium Azide
Isotype	IgG
Purity	Immunogen affinity purified
Buffer	PBS

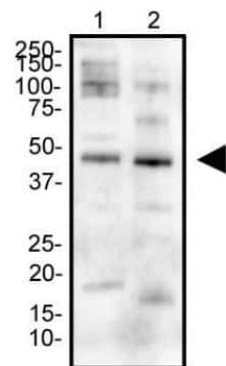
Product Description	
Host	Rabbit
Gene ID	2057
Gene Symbol	EPOR
Species	Human, Mouse, Rat
Reactivity Notes	Use in Rat reported in scientific literature (PMID:34552120).
Specificity/Sensitivity	EpoR (A364) detects endogenous levels of EpoR protein.
Immunogen	A synthetic peptide made to an internal portion of the human EPO receptor protein (between residues 300-400) [UniProt P19235]

Product Application Details	
Applications	Western Blot, Simple Western, Immunocytochemistry/Immunofluorescence, Immunohistochemistry, Immunohistochemistry-Paraffin
Recommended Dilutions	Western Blot 2.0 ug/ml, Simple Western 1:10, Immunohistochemistry 1:50 - 1:100, Immunocytochemistry/Immunofluorescence 1:50 - 1:100, Immunohistochemistry-Paraffin 1:50-1:100
Application Notes	In Simple Western only 10 - 15 uL of the recommended dilution is used per data point. Separated by Size.

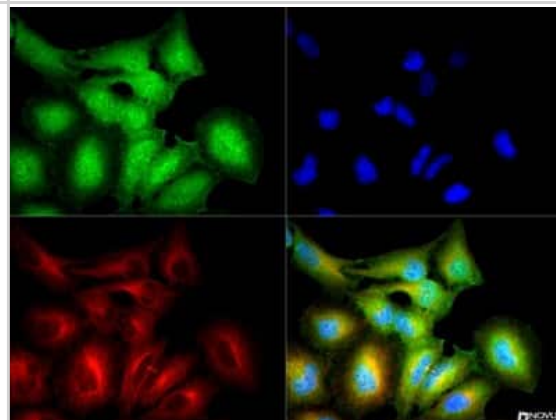


Images

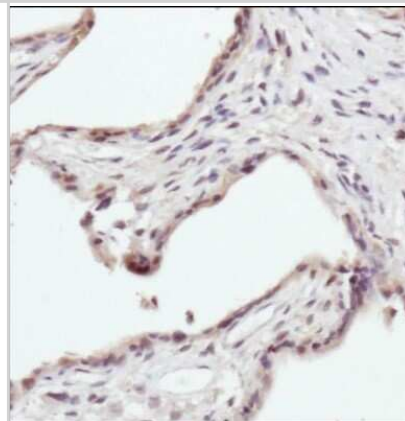
Western Blot: Erythropoietin R Antibody [NBP1-19388] - Western blot analysis of K562 (left) and SH-SY5Y (right) cell lysate using EPO receptor antibody at a concentration of 2 ug/ml.



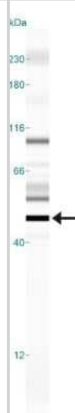
Immunocytochemistry/Immunofluorescence: Erythropoietin R Antibody [NBP1-19388] - EPO receptor antibody was tested in HeLa cells with DyLight 488 (green). Nuclei and alpha-tubulin were counterstained with DAPI (blue) and Dylight 550 (red).



Immunohistochemistry-Paraffin: Erythropoietin R Antibody [NBP1-19388] - IHC analysis of the EPO receptor in human placenta.



Simple Western: Erythropoietin R Antibody [NBP1-19388] - Simple Western lane view shows a specific band for EPO Receptor in 0.5 mg/ml of Human Brain lysate. This experiment was performed under reducing conditions using the 12-230 kDa separation system.



Publications

Zhang L, Liu X, Wei Q et al. Arginine attenuates chronic mountain sickness in rats via microRNA-144-5p Mammalian genome : official journal of the International Mammalian Genome Society 2023-03-01 [PMID: 36763178] (IHC)

Hu SQ, Liu DL, Li CR et al. Wuzi-Yanzong prescription alleviates spermatogenesis disorder induced by heat stress dependent on Akt, NF- kappaB signaling pathway Scientific reports 2021-09-22 [PMID: 34552120] (Rat)

Vaughan OR, Thompson F, Lorca RA EFFECT OF HIGH ALTITUDE ON HUMAN PLACENTAL AMINO ACID TRANSPORT J. Appl. Physiol. 2019-12-05 [PMID: 31804891]

Tanaka T, Murata H, Miyazaki R, et al. Human recombinant erythropoietin improves motor function in rats with spinal cord compression-induced cervical myelopathy PLoS ONE 2019-12-10 [PMID: 31821342] (IF/IHC, Rat)



Procedures

Western blot protocol for EPO receptor antibody (NBP1-19388)

Erythropoietin R Antibody: https://www.novusbio.com/products/erythropoietin-r-antibody_nbp1-19388

Western Blot Protocol

1. Perform SDS-PAGE on samples to be analyzed, loading 25 ug of total protein per lane.
2. Transfer proteins to membrane according to the instructions provided by the manufacturer of the membrane and transfer apparatus.
3. Stain according to standard Ponceau S procedure (or similar product) to assess transfer success, and mark molecular weight standards where appropriate.
4. Rinse the blot.
5. Block the membrane using standard blocking buffer for at least 1 hour.
6. Wash the membrane in wash buffer three times for 10 minutes each.
7. Dilute anti-HDAC5 primary antibody in blocking buffer and incubate 1 hour at room temperature.
8. Wash the membrane in wash buffer three times for 10 minutes each.
9. Apply the diluted HRP conjugated secondary antibody in blocking buffer (as per manufacturers instructions) and incubate 1 hour at room temperature.
10. Wash the blot in wash buffer three times for 10 minutes each (this step can be repeated as required to reduce background).
11. Apply the detection reagent of choice in accordance with the manufacturers instructions.

Note: Tween-20 can be added to the blocking or antibody dilution buffer at a final concentration of 0.05-0.2%.

Immunohistochemistry-Paraffin protocol for EPO receptor antibody (NBP1-19388)

Erythropoietin R Antibody: https://www.novusbio.com/products/erythropoietin-r-antibody_nbp1-19388

Immunohistochemistry-Paraffin Embedded Sections

Antigen Unmasking:

Bring slides to a boil in 10 mM sodium citrate buffer (pH 6.0) then maintain at a sub-boiling temperature for 10 minutes. Cool slides on bench-top for 30 minutes.

Staining:

1. Wash sections in deionized water three times for 5 minutes each.
2. Wash sections in wash buffer for 5 minutes.
3. Block each section with 100-400 ul blocking solution for 1 hour at room temperature.
4. Remove blocking solution and add 100-400 ul diluted primary antibody. Incubate overnight at 4 C.
5. Remove antibody solution and wash sections in wash buffer three times for 5 minutes each.
6. Add 100-400 ul biotinylated diluted secondary antibody. Incubate 30 minutes at room temperature.
7. Remove secondary antibody solution and wash sections three times with wash buffer for 5 minutes each.
8. Add 100-400 ul Streptavidin-HRP reagent to each section and incubate for 30 minutes at room temperature.
9. Wash sections three times in wash buffer for 5 minutes each.
10. Add 100-400 ul DAB substrate to each section and monitor staining closely.
11. As soon as the sections develop, immerse slides in deionized water.
12. Counterstain sections in hematoxylin.
13. Wash sections in deionized water two times for 5 minutes each.
14. Dehydrate sections.
15. Mount coverslips.



Immunocytochemistry/Immunofluorescence protocol for EPO receptor antibody (NBP1-19388)Erythropoietin R Antibody: https://www.novusbio.com/products/erythropoietin-r-antibody_nbp1-19388

Immunocytochemistry Protocol

Culture cells to appropriate density in 35 mm culture dishes or 6-well plates.

1. Remove culture medium and add 10% formalin to the dish. Fix at room temperature for 30 minutes.
2. Remove the formalin and add ice cold methanol. Incubate for 5-10 minutes.
3. Remove methanol and add washing solution (i.e. PBS). Be sure to not let the specimen dry out. Wash three times for 10 minutes.
4. To block nonspecific antibody binding incubate in 10% normal goat serum from 1 hour to overnight at room temperature.
5. Add primary antibody at appropriate dilution and incubate at room temperature from 2 hours to overnight at room temperature.
6. Remove primary antibody and replace with washing solution. Wash three times for 10 minutes.
7. Add secondary antibody at appropriate dilution. Incubate for 1 hour at room temperature.
8. Remove antibody and replace with wash solution, then wash for 10 minutes. Add Hoechst 33258 to wash solution at 1:25,000 and incubate for 10 minutes. Wash a third time for 10 minutes.
9. Cells can be viewed directly after washing. The plates can also be stored in PBS containing Azide covered in Parafilm (TM). Cells can also be cover-slipped using Fluoromount, with appropriate sealing.

*The above information is only intended as a guide. The researcher should determine what protocol best meets their needs. Please follow safe laboratory procedures.





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Products Related to NBP1-19388

HAF008	Goat anti-Rabbit IgG Secondary Antibody [HRP]
NB7160	Goat anti-Rabbit IgG (H+L) Secondary Antibody [HRP]
NBP2-24891	Rabbit IgG Isotype Control
NBP1-19388B	Erythropoietin R Antibody [Biotin]

Limitations

This product is for research use only and is not approved for use in humans or in clinical diagnosis. Primary Antibodies are guaranteed for 1 year from date of receipt.

For more information on our 100% guarantee, please visit www.novusbio.com/guarantee

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