Product Datasheet

RUNX2/CBFA1 Antibody - BSA Free NBP1-77461

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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NBP1-77461

RUNX2/CBFA1 Antibody - BSA Free

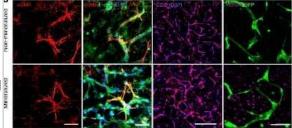
RUNX2/CBFA1 Antibody - BSA Free	
Product Information	
Unit Size	0.1 ml
Concentration	1 mg/ml
Storage	Store at 4C short term. Aliquot and store at -20C long term. Avoid freeze-thaw cycles.
Clonality	Polyclonal
Preservative	0.05% Sodium Azide
Isotype	IgG
Purity	Immunogen affinity purified
Buffer	PBS and 30% Glycerol
Target Molecular Weight	56.6 kDa
Product Description	
Host	Rabbit
Gene ID	860
Gene Symbol	RUNX2
Species	Human, Mouse
Immunogen	This RUNX2/CBFA1 Antibody was prepared from a synthetic peptide made to an internal region of the human RUNX2 protein (within residues 300-375). [Swiss-Prot Q13950]
Product Application Details	
Applications	Western Blot, Immunocytochemistry/ Immunofluorescence, Immunohistochemistry, Immunohistochemistry-Paraffin
Recommended Dilutions	Western Blot reported in scientific literature (PMID 29208768), Immunohistochemistry 1:200, Immunocytochemistry/ Immunofluorescence 1:100, Immunohistochemistry-Paraffin 1:200
Application Notes	Prior to immunostaining paraffin tissues antigen retrieval with sodium citrate

buffer (pH 6.0) is recommended.

Images

Immunohistochemistry-Paraffin: RUNX2/CBFA1 Antibody - BSA Free [NBP1-77461] - RUNX2/CBFA1 Antibody [NBP1-77461] - Vascularization of mineralized cell-laden collagen and interaction with prostate cancer cells. HUVECs formed endothelial networks that were supported by SMA-expressing hMSCs (scale bar: 50m) and were also positive for CD31 (scale bar: 400m). The remainder of hMSCs expressed RUNX2 as a marker for osteogenic differentiation (scale bar: 50m). Rapid fabrication of vascularized and innervated cell-laden bone models with biomimetic intrafibrillar collagen mineralization. Image collected and cropped by CiteAb from the following publication

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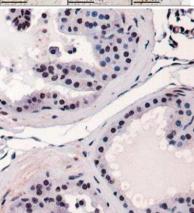


Immunocytochemistry/Immunofluorescence: RUNX2/CBFA1 Antibody [NBP1-77461] - HeLa cells were fixed in 4% paraformaldehyde for 10 minutes and permeabilized in 0.5% Triton X-100 in PBS for 5 minutes. The cells were incubated with RUNX2/CBFA1 Antibody (NBP1-77461) at 1 ug/ml overnight at 4C and detected with an anti-rabbit Dylight 488 (Green) at a 1:1000 dilution for 60 minutes. Nuclei were counterstained with DAPI (Blue). Cells were imaged using a 100X objective and digitally deconvolved.



Immunohistochemistry-Paraffin: RUNX2/CBFA1 Antibody - BSA Free [NBP1-77461] - RUNX2/CBFA1 Antibody [NBP1-77461] - Characterization of TLC cultured in chondrogenic pellets. Gene expression of TLC cultured four weeks in chondrogenic pellets or in cell monolayer in control medium: levels of expression of osteoblast markers (1), chondrocyte markers (2) and tenocyte markers (3) (N = 3) assessed by qPCR.* =0.05. Rotator Cuff Tenocytes Differentiate into Hypertrophic Chondrocyte-Like Cells to Produce Calcium Deposits in an Alkaline Phosphatase-Dependent Manner. Image collected and cropped by CiteAb from the following publication (//pubmed.ncbi.nlm.nih.gov/31561454/) licensed under a CC-BY license.

Immunohistochemistry: RUNX2/CBFA1 Antibody [NBP1-77461] - Staining of RUNX2 in mouse prostate using DAB with hematoxylin counterstain.



Immunohistochemistry: RUNX2/CBFA1 Antibody - BSA Free [NBP1-77461] - Histological characterization of rotator cuff calcifications (N = 5). (A) Calcific deposits assessed by micro-computed tomography. (B) HE (hematoxylin & eosin) staining of decalcified samples (1 image for each patient). (C) Von Kossa staining on not decalcified samples & characterization of the fibrocartilaginous area with alcian blue staining, SOFG (Safranin O/Fast Green) staining, & immunohistochemical staining for COL2 (brown) with Gill hematoxylin counterstain. (D,E) Representative immunohistochemical staining for ENPP1 (ectonucleotide pyrophosphatase/phosphodiesterase 1), TNAP (tissue non-specific alkaline phosphatase), Runx2, Sox9, Col10 & CD31 (brown). TF: tendon fibers; CD: calcium deposits, FM: fibrocartilaginous metaplasia; OM: osseous metaplasia; CT: connective tissue; V: vessels. Scale bar = 200 µM. Image collected & cropped by CiteAb from the following publication (https://pubmed.ncbi.nlm.nih.gov/31561454), licensed under a CC-BY license. Not internally tested by Novus Biologicals.



Publications

Sousa MGDC, Balbinot GS, Subbiah R et al. In vitro development and optimization of cell-laden injectable bioprinted gelatin methacryloyl (GelMA) microgels mineralized on the nanoscale bioRxiv: the preprint server for biology 2023-10-12 [PMID: 37873385] (Immunocytochemistry/ Immunofluorescence, Mouse)

Chan, B;Glogauer, M;Wang, Y;Wrana, J;Chan, K;Beier, F;Bali, S;Hinz, B;Parreno, J;Ashraf, S;Kandel, R; Adseverin, an actin-binding protein, modulates hypertrophic chondrocyte differentiation and osteoarthritis progression Science advances 2023-08-04 [PMID: 37540756] (In vivo assay)

Poudel SB, Ruff RR, Yildirim G et al. Excess growth hormone triggers inflammation-associated arthropathy, subchondral bone loss, and arthralgia The American journal of pathology 2023-03-02 [PMID: 36870529] (IHC-P, Mouse)

Subbiah R, Lin EY, Athirasala A et al. Engineering of an osteoinductive and growth factor-free injectable bone-like microgel for bone regeneration Advanced healthcare materials 2023-02-20 [PMID: 36808718]

Zanut A, Li R, Deng R et al. A polymer canvas with the stiffness of the bone matrix to study and control mesenchymal stem cell response Advanced healthcare materials 2022-12-24 [PMID: 36565136]

Ciavarella C, Motta I, Vasuri F et al. Involvement of miR-30a-5p and miR-30d in Endothelial to Mesenchymal Transition and Early Osteogenic Commitment under Inflammatory Stress in HUVEC Biomolecules 2021-02-05 [PMID: 33562690] (ICC/IF, Human)

Serjeant M, Moon PM, Quinonez D, et al. The Role of Panx3 in Age-Associated and Injury-Induced Intervertebral Disc Degeneration International journal of molecular sciences 2021-01-22 [PMID: 33499145] (IF/IHC, Mouse)

Cahill SV, Kwon HK, Back J et al. Locally-delivered Adjuvant Biofilm-penetrating Antibiotics Rescue Impaired Endochondral Fracture Healing Caused by MRSA Infection Journal of orthopaedic research: official publication of the Orthopaedic Research Society 2020-12-18 [PMID: 33336805]

Chen PY, Qin L, Li G et al. Smooth Muscle Cell Reprogramming in Aortic Aneurysms Cell Stem Cell 2020-04-02 [PMID: 32243809] (Mouse)

Darrieutort-Laffite C, Arnolfo P, Garraud T et al. Rotator Cuff Tenocytes Differentiate into Hypertrophic Chondrocyte-Like Cells to Produce Calcium Deposits in an Alkaline Phosphatase-Dependent Manner J Clin Med [PMID: 31561454] (IF/IHC)

Thrivikraman G, Athirasala A, Gordon R et al. Rapid fabrication of vascularized and innervated cell-laden bone models with biomimetic intrafibrillar collagen mineralization Nat Commun 2019-08-06 [PMID: 31388010] (ICC/IF, Human)

Priante G, Ceol M, Gianesello L et al. Human proximal tubular cells can form calcium phosphate deposits in osteogenic culture: role of cell death and osteoblast-like transdifferentiation. Cell Death Discov 2019-01-28 [PMID: 30701089] (WB, Human)

More publications at http://www.novusbio.com/NBP1-77461



Procedures

Immunohistochemistry-Paraffin protocol for RUNX2 Antibody (NBP1-77461)

RUNX2/CBFA1 Antibody:

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 4C.
- 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.

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

Immunocytochemistry/Immunofluorescence Protocol for RUNX2 Antibody (NBP1-77461)

RUNX2/CBFA1 Antibody:

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 1:25,0000 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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HAF008 Goat anti-Rabbit IgG Secondary Antibody [HRP]

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NBP2-24891 Rabbit IgG Isotype Control

NBP1-77461AF647 RUNX2/CBFA1 Antibody [Alexa Fluor® 647]

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