BOSTER BIOLOGICAL TECHNOLOGY
3942 B Valley Ave, Pleasanton, CA 94566
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Anti-SMARCD3 Antibody

Catalog Number: A06464-1

About SMARCD3

SMARCD3 is a component of the SWI/SNF (mating type switching/sucrose non-fermenting) chromatin remodeling complex, which evolutionarily conserved from yeast to human, and all complexes contain a core set of conserved components, including BRG1/Brm-associated factors (BAFs) and a DNA-dependent SWI2/SNF2-like ATPase, which enables chromatin remodeling. SMARCD3 isoforms bind to several nuclear receptors and transcription factors of various families. SMARCD3 proteins interact in a ligand-independent manner with peroxisome proliferator- activated receptor gamma and enhance its transcriptional activity.

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Overview

Product Name	Anti-SMARCD3 Antibody
Reactive Species	Human, Mouse, Rat
Description	Boster Bio Anti-SMARCD3 Antibody catalog # A06464-1. Tested in WB, IHC, IF, ELISA applications. This antibody reacts with Human, Mouse, Rat.
Application	ELISA, IF, IHC, WB
Clonality	Polyclonal
Formulation	500 ug/ml antibody with PBS, 0.02% NaN3, 1 mg stabilizing protein and 50% glycerol *This antibody is supplied in a stabilized formulation. Compatibility with conjugation reactions depends on the chemistry of the conjugation method used. For conjugation methods that are not compatible with the stabilizing components present in this formulation, a carrier-free antibody format is required.
Storage Instructions	12 months from date of receipt□-20°C as supplied. 6 months 2 to 8°C after reconstitution. Avoid repeated freezing and thawing.
Host	Rabbit
Uniprot ID	Q6STE5

Technical Details

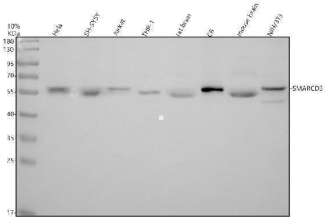
Immunogen	E.coli-derived human SMARCD3 recombinant protein (Position: P3-A85).

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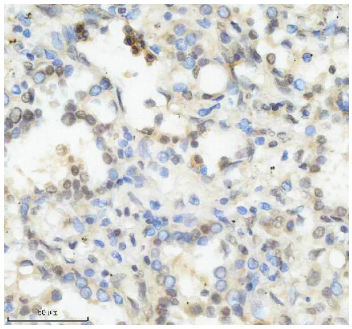
Form	Liquid
Concentration	500 ug/ml
Purification	Immunogen affinity purified.
Suggested Dilutions	Western blot, 1:500-2000 Immunohistochemistry, 1:50-400 Immunofluorescence, 1:50-400 ELISA, 1:100-1000

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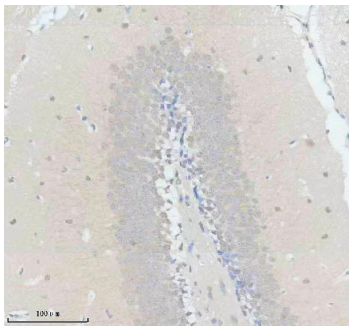
Anti-SMARCD3 Antibody (A06464-1) Images



Western blot analysis of SMARCD3 using anti-SMARCD3 antibody (A06464-1). Electrophoresis was performed on a 10% SDS-PAGE gel at 80V (Stacking gel) / 120V (Resolving gel) for 2 hours. The sample well of each lane was loaded with 30 ug of sample under reducing conditions. Lane 1: human Hela whole cell lysates, Lane 2: human SH-SY5Y whole cell lysates, Lane 3: human Jurkat whole cell lysates, Lane 4: human THP-1 whole cell lysates, Lane 5: rat brain tissue lysates, Lane 6: rat C6 whole cell lysates, Lane 7: mouse brain tissue lysates, Lane 8: mouse NIH/3T3 whole cell lysates.After electrophoresis, proteins were transferred to a nitrocellulose membrane at 150 mA for 50-90 minutes. Blocked the membrane with 5% non-fat milk/TBS for 1.5 hour at RT. The membrane was incubated with rabbit anti-SMARCD3 antigen affinity purified polyclonal antibody (A06464-1) at 1:1000 overnight at 4°C, then washed with TBS-0.1%Tween 3 times with 5 minutes each and probed with a goat anti-rabbit IgG-HRP secondary antibody at a dilution of 1:5000 for 1.5 hour at RT. The signal is developed using an ECL Plus Western Blotting Substrate (Catalog # AR1196-200) with Tanon 5200 system. A specific band was detected for SMARCD3 at approximately 60 kDa. The expected band size for SMARCD3 is at 55 kDa.



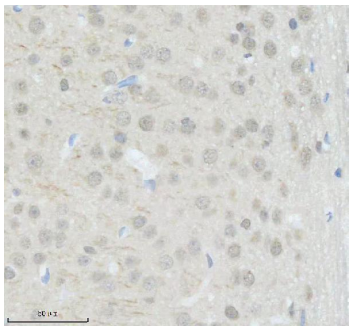
IHC analysis of SMARCD3 using anti-SMARCD3 antibody (A06464-1). SMARCD3 was detected in a paraffin-embedded section of human lung cancer tissue. Heat mediated antigen retrieval was performed in EDTA buffer (pH 8.0, epitope retrieval solution). The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 1:200 rabbit anti-SMARCD3 Antibody (A06464-1) overnight at 4°C. Peroxidase Conjugated Goat Anti-rabbit IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using HRP Conjugated Rabbit IgG Super Vision Assay Kit (Catalog # SV0002) with DAB as the chromogen.



IHC analysis of SMARCD3 using anti-SMARCD3 antibody (A06464-1). SMARCD3 was detected in a paraffin-embedded section of mouse brain tissue. Heat mediated antigen retrieval was performed in EDTA buffer (pH 8.0, epitope retrieval solution). The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 1:200 rabbit anti-SMARCD3 Antibody (A06464-1) overnight at 4°C. Peroxidase Conjugated Goat Anti-rabbit IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using HRP Conjugated Rabbit IgG Super Vision Assay Kit (Catalog # SV0002) with DAB as the chromogen.

IHC analysis of SMARCD3 using anti-SMARCD3 antibody

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Anti-SMARCD3 Antibody

For Research Use Only. Not for use in diagnostic procedures.