

Anti-Astrin/Deepest/SPAG5 Antibody Picoband™

Catalog Number: A07062-2

About SPAG5

Sperm-associated antigen 5 is a protein that in humans is encoded by the SPAG5 gene. This gene encodes a protein associated with the mitotic spindle apparatus. The encoded protein may be involved in the functional and dynamic regulation of mitotic spindles.

Overview

Product Name	Anti-Astrin/Deepest/SPAG5 Antibody Picoband™
Reactive Species	Human
Description	Boster Bio Anti-Astrin/Deepest/SPAG5 Antibody Picoband™ catalog # A07062-2. Tested in ELISA, Flow Cytometry, IF, IHC-P, ICC, WB applications. This antibody reacts with Human.
Application	ELISA, Flow Cytometry, IF, IHC-P, ICC, WB
Clonality	Polyclonal
Formulation	Each vial contains 4mg Trehalose, 0.9mg NaCl, 0.2mg Na ₂ HPO ₄ .
Storage Instructions	Store at -20°C for one year from date of receipt. After reconstitution, at 4°C for one month. It can also be aliquotted and stored frozen at -20°C for six months. Avoid repeated freeze-thaw cycles.
Host	Rabbit
Uniprot ID	Q96R06

Technical Details

Immunogen	E.coli-derived human Astrin/Deepest/SPAG5 recombinant protein (Position: E738-S1193).
Recommended Detection Systems	Boster recommends Enhanced Chemiluminescent Kit with anti-Rabbit IgG (EK1002) for Western blot, and HRP Conjugated anti-Rabbit IgG Super Vision Assay Kit (SV0002-1) for IHC(P) and ICC.
Form	Lyophilized
Concentration	Adding 0.2 ml of distilled water will yield a concentration of 500 µg/ml.
Purification	Immunogen affinity purified.
Suggested Dilutions	Dilute the sample so that the expected range of concentrations fall within the detection range of this

kit.

If the expected range of concentration is unknown, a pilot test should be conducted to decide the optimal dilution ratio for your samples.

Some PubMed article(s) citing the expression level of this target are as follows:

Boster Bio's internal QC testing used Western blot, 0.25-0.5 μ g/ml, Human

Immunohistochemistry (Paraffin-embedded Section), 2-5 μ g/ml, Human

Immunocytochemistry/Immunofluorescence, 5 μ g/ml, Human

Flow Cytometry, 1-3 μ g/1x10⁶ cells, Human

Direct ELISA, 0.1-0.5 μ g/ml, Human

For protocols please visit <https://www.bosterbio.com/protocol-and-troubleshooting/>

Anti-Astrin/Deepest/SPAG5 Antibody Picoband™ (A07062-2) Images

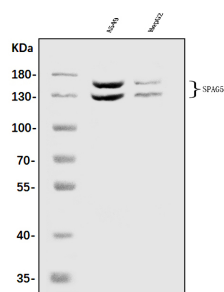


Figure 1. Western blot analysis of Astrin/Deepest/SPAG5 using anti-Astrin/Deepest/SPAG5 antibody (A07062-2). Electrophoresis was performed on a 5-20% SDS-PAGE gel at 70V (Stacking gel) / 90V (Resolving gel) for 2-3 hours. The sample well of each lane was loaded with 30 μ g of sample under reducing conditions.

Lane 1: human A549 whole cell lysates,

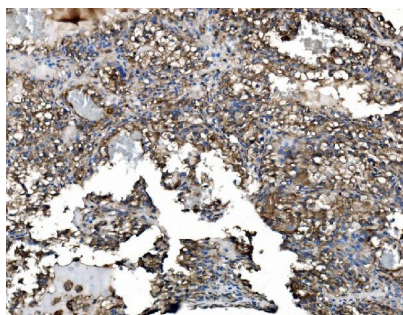
Lane 2: human HepG2 whole cell lysates.

After Electrophoresis, proteins were transferred to a Nitrocellulose membrane at 150mA 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-Astrin/Deepest/SPAG5 antigen affinity purified polyclonal antibody (Catalog # A07062-2) at 0.5 μ g/mL 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 Enhanced Chemiluminescent detection (ECL) kit (Catalog # EK1002) with Tanon 5200 system. A specific band was detected for Astrin/Deepest/SPAG5 at approximately 130KD, 150KD. The expected band size for Astrin/Deepest/SPAG5 is at 130KD, 150KD.

Figure 2. IHC analysis of Astrin/Deepest/SPAG5 using anti-Astrin/Deepest/SPAG5 antibody (A07062-2).

Astrin/Deepest/SPAG5 was detected in paraffin-embedded section of human renal clear cell carcinoma tissue. Heat mediated antigen retrieval was performed in EDTA buffer (pH8.0, epitope retrieval solution). The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 2 μ g/ml rabbit anti-Astrin/Deepest/SPAG5 Antibody (A07062-2) overnight at 4°C. Biotinylated goat anti-



rabbit IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using Streptavidin-Biotin-Complex (SABC) (Catalog # SA1022) with DAB as the chromogen.

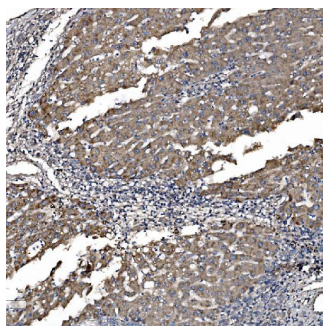


Figure 3. IHC analysis of Astrin/Deepest/SPAG5 using anti-Astrin/Deepest/SPAG5 antibody (A07062-2). Astrin/Deepest/SPAG5 was detected in paraffin-embedded section of human liver cancer tissue. Heat mediated antigen retrieval was performed in EDTA buffer (pH8.0, epitope retrieval solution). The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 2ug/ml rabbit anti-Astrin/Deepest/SPAG5 Antibody (A07062-2) overnight at 4°C. Biotinylated goat anti-rabbit IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using Streptavidin-Biotin-Complex (SABC) (Catalog # SA1022) with DAB as the chromogen.

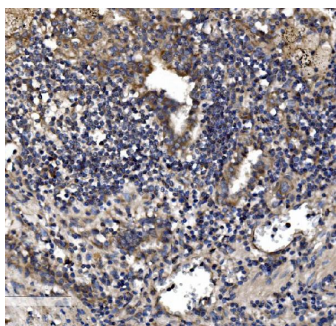


Figure 4. IHC analysis of Astrin/Deepest/SPAG5 using anti-Astrin/Deepest/SPAG5 antibody (A07062-2). Astrin/Deepest/SPAG5 was detected in paraffin-embedded section of human lung cancer tissue. Heat mediated antigen retrieval was performed in EDTA buffer (pH8.0, epitope retrieval solution). The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 2ug/ml rabbit anti-Astrin/Deepest/SPAG5 Antibody (A07062-2) overnight at 4°C. Biotinylated goat anti-rabbit IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using Streptavidin-Biotin-Complex (SABC) (Catalog # SA1022) with DAB as the chromogen.

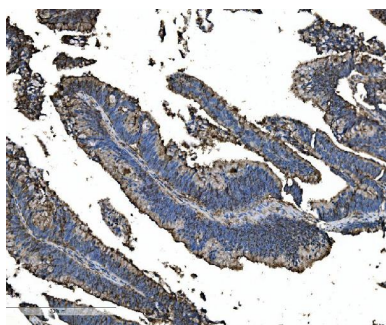


Figure 5. IHC analysis of Astrin/Deepest/SPAG5 using anti-Astrin/Deepest/SPAG5 antibody (A07062-2). Astrin/Deepest/SPAG5 was detected in paraffin-embedded section of human rectal cancer tissue. Heat mediated antigen retrieval was performed in EDTA buffer (pH8.0, epitope retrieval solution). The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 2ug/ml rabbit anti-Astrin/Deepest/SPAG5 Antibody (A07062-2) overnight at 4°C. Biotinylated goat anti-rabbit IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using Streptavidin-Biotin-Complex (SABC) (Catalog # SA1022) with DAB as the chromogen.

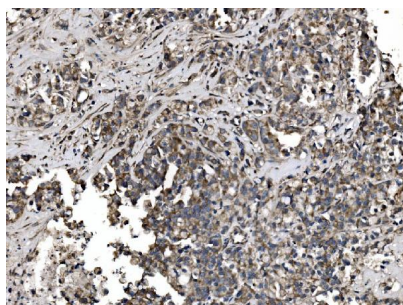


Figure 6. IHC analysis of Astrin/Deepest/SPAG5 using anti-Astrin/Deepest/SPAG5 antibody (A07062-2). Astrin/Deepest/SPAG5 was detected in paraffin-embedded section of human gastric cancer tissue. Heat mediated antigen retrieval was performed in EDTA buffer (pH8.0, epitope retrieval solution). The tissue section was blocked with 10% goat serum. The tissue section was then incubated with 2ug/ml rabbit anti-Astrin/Deepest/SPAG5 Antibody (A07062-2) overnight at 4°C. Biotinylated goat anti-rabbit IgG was used as secondary antibody and incubated for 30 minutes at 37°C. The tissue section was developed using Streptavidin-Biotin-Complex (SABC) (Catalog # SA1022) with DAB as the chromogen.

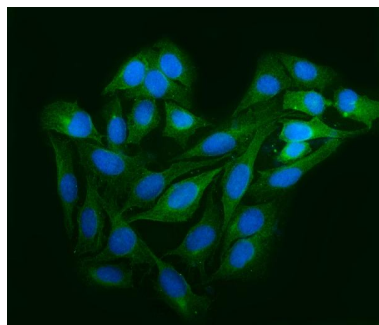


Figure 7. IF analysis of Astrin/Deepest/SPAG5 using anti-Astrin/Deepest/SPAG5 antibody (A07062-2). Astrin/Deepest/SPAG5 was detected in immunocytochemical section of U20S cells. Enzyme antigen retrieval was performed using IHC enzyme antigen retrieval reagent (AR0022) for 15 mins. The cells were blocked with 10% goat serum. And then incubated with 5ug/mL rabbit anti-Astrin/Deepest/SPAG5 Antibody (A07062-2) overnight at 4°C. DyLight®488 Conjugated Goat Anti-Rabbit IgG (BA1127) was used as secondary antibody at 1:100 dilution and incubated for 30 minutes at 37°C. The section was counterstained with DAPI. Visualize using a fluorescence microscope and filter sets appropriate for the label used.

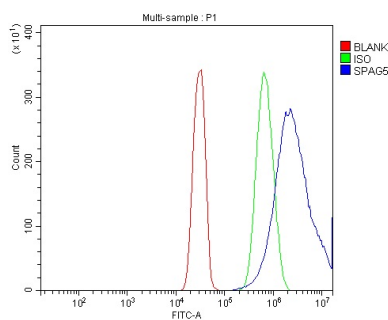


Figure 8. Flow Cytometry analysis of SiHa cells using anti-Astrin/Deepest/SPAG5 antibody (A07062-2). Overlay histogram showing SiHa cells stained with A07062-2 (Blue line). The cells were blocked with 10% normal goat serum. And then incubated with rabbit anti-Astrin/Deepest/SPAG5 Antibody (A07062-2, 1ug/1x10⁶ cells) for 30 min at 20°C. DyLight®488 conjugated goat anti-rabbit IgG (BA1127, 5-10ug/1x10⁶ cells) was used as secondary antibody for 30 minutes at 20°C. Isotype control antibody (Green line) was rabbit IgG (1ug/1x10⁶) used under the same conditions. Unlabelled sample (Red line) was also used as a control.

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