

# Product Datasheet

## FUSIP1 Antibody NB110-93598

Unit Size: 0.1 ml

Store at 4C. Do not freeze.

[www.novusbio.com](http://www.novusbio.com)



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**NB110-93598****FUSIP1 Antibody**

| <b>Product Information</b>     |                                       |
|--------------------------------|---------------------------------------|
| <b>Unit Size</b>               | 0.1 ml                                |
| <b>Concentration</b>           | 1 mg/ml                               |
| <b>Storage</b>                 | Store at 4C. Do not freeze.           |
| <b>Clonality</b>               | Polyclonal                            |
| <b>Preservative</b>            | 0.1% Sodium Azide                     |
| <b>Isotype</b>                 | IgG                                   |
| <b>Purity</b>                  | Immunogen affinity purified           |
| <b>Buffer</b>                  | Tris-Citrate/Phosphate (pH 7.0 - 8.0) |
| <b>Target Molecular Weight</b> | 22 kDa                                |

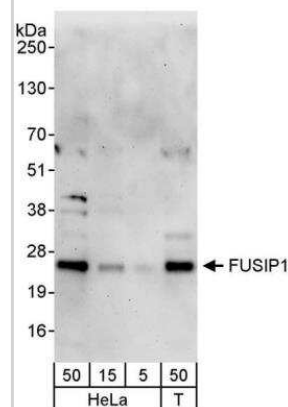
| <b>Product Description</b>     |   |
|--------------------------------|---|
| <b>Host</b>                    | Rabbit  |
| <b>Gene ID</b>                 | 10772   |
| <b>Gene Symbol</b>             | SRSF10  |
| <b>Species</b>                 | Human   |
| <b>Specificity/Sensitivity</b> | This is specific for isoform 3 of the FUSIP1 protein.   |
| <b>Immunogen</b>               | Synthetic peptide made to an internal portion of human FUSIP1 (within residues 100-200). [Swiss-Prot# O75494] |

| <b>Product Application Details</b> |   |
|------------------------------------|---|
| <b>Applications</b>                | Western Blot, Immunocytochemistry/Immunofluorescence, Immunohistochemistry-Paraffin   |
| <b>Recommended Dilutions</b>       | Western Blot 1:1000 -1:5000, Immunocytochemistry/Immunofluorescence 1:1000 - 1:2000, Immunohistochemistry-Paraffin  |
| <b>Application Notes</b>           | This FUSIP1 antibody is useful for Immunocytochemistry/Immunofluorescence and Western blot, where a band is seen ~22 kDa. The observed molecular weight of the protein may vary from the listed predicted molecular weight due to post translational modifications, post translation cleavages, relative charges, and other experimental factors. |

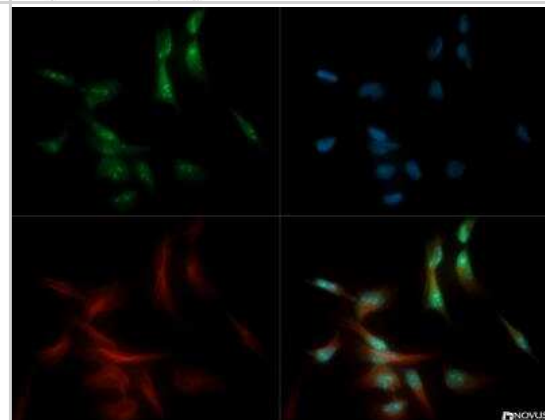


## Images

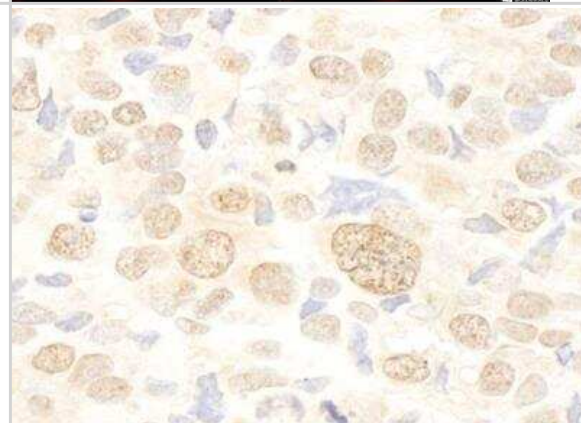
**Western Blot: FUSIP1 Antibody [NB110-93598]** - Detection of human FUSIP1 by western blot. Samples: Whole cell lysate from HeLa (5, 15 and 50 ug) and HEK293T (T; 50 ug) cells. Antibody: Affinity purified rabbit anti-FUSIP1 antibody NB110-93598 used for WB at 0.4 ug/ml. Detection: Chemiluminescence with an exposure time of 3 minutes.



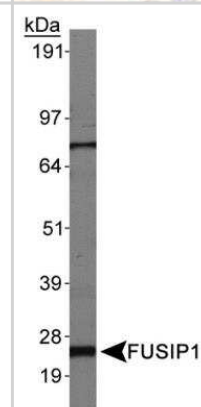
**Immunocytochemistry/Immunofluorescence: FUSIP1 Antibody [NB110-93598]** - FUSIP1 antibody was tested in Hela cells with DyLight 488 (green). Nuclei and alpha-tubulin were counterstained with DAPI (blue) and Dylight 550 (red).



**Immunohistochemistry-Paraffin: FUSIP1 Antibody [NB110-93598]** - Detection of human FUSIP1 by immunohistochemistry. Sample: FFPE section of human breast carcinoma. Antibody: Affinity purified rabbit anti-FUSIP1 (NB110-93598) used at a dilution of 1:1,000 (1ug/ml). Detection: DAB



**Western Blot: FUSIP1 Antibody [NB110-93598]** - Detection of FUSIP1, Isoform 3, in HepG2 whole cell lysates using NB110-93598.



## Publications

Fan Y, Zhang Y, Zhao H et al. lncR-GAS5 upregulates the splicing factor SRSF10 to impair endothelial autophagy, leading to atherogenesis *Frontiers of medicine* 2023-04-01 [PMID: 36645633]

Liu W, Lu X, Zhao ZH et al. SRSF10 is essential for progenitor spermatogonia expansion by regulating alternative splicing *eLife* 2022-11-10 [PMID: 36355419] (WB, Mouse)



## Procedures

### Western Blot protocol for FUSIP1 Antibody (NB110-93598)

FUSIP1 Antibody: [https://www.novusbio.com/products/fusip1-antibody\\_nb110-93598](https://www.novusbio.com/products/fusip1-antibody_nb110-93598)

#### Western Blot Protocol

1. Perform SDS-PAGE (4-12% MOPS) on samples to be analyzed, loading 30 ug of total protein per lane.
2. Transfer proteins to Nitrocellulose according to the instructions provided by the manufacturer of the transfer apparatus.
3. Rinse membrane with dH<sub>2</sub>O and then stain the blot using Ponceau S for 1-2 minutes to access the transfer of proteins onto the nitrocellulose membrane. Rinse the blot in water to remove excess stain and mark the lane locations and locations of molecular weight markers using a pencil.
4. Rinse the blot in TBS for approximately 5 minutes.
5. Block the membrane using 5% NFDM + 1% BSA in TBS + Tween, 1 hour at RT.
6. Rinse the membrane in dH<sub>2</sub>O and then wash the membrane in wash buffer [TBS + 0.1% Tween] 3 times for 10 minutes each.
7. Dilute the rabbit anti-FUSIP1 primary antibody (NB 110-93598) in blocking buffer and incubate 1 hour at room temperature.
8. Rinse the membrane in dH<sub>2</sub>O and then wash the membrane in wash buffer [TBS + 0.1% Tween] 3 times for 10 minutes each.
9. Apply the diluted rabbit-IgG 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 [TBS + 0.1% Tween] 3 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 (Pierce ECL).

Note: Tween-20 can be added to the blocking or antibody dilution buffer at a final concentration of 0.05-0.2%, provided it does not interfere with antibody-antigen binding.

### Immunocytochemistry/Immunofluorescence protocol for FUSIP1 Antibody (NB110-93598)

FUSIP1 Antibody: [https://www.novusbio.com/products/fusip1-antibody\\_nb110-93598](https://www.novusbio.com/products/fusip1-antibody_nb110-93598)

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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### **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](http://www.novusbio.com/guarantee)

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