# **Product Datasheet**

# DUSP22 Antibody - BSA Free NLS6809

Unit Size: 0.05 ml

Keep as concentrated solution. Aliquot and store at -20C or below. Avoid multiple freeze-thaw cycles.

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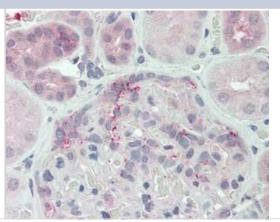
#### NLS6809

DUSP22 Antibody - BSA Free

Product Information	
Unit Size	0.05 ml
Concentration	1.0 mg/ml
Storage	Keep as concentrated solution. Aliquot and store at -20C or below. Avoid multiple freeze-thaw cycles.
Clonality	Polyclonal
Preservative	0.1% Sodium Azide
Isotype	IgG
Purity	Immunogen affinity purified
Buffer	PBS
Product Description	
Description	Product can be stored undiluted at 4C for up to 1 month.
Host	Rabbit
Gene ID	56940
Gene Symbol	DUSP22
Species	Human, Bovine, Equine, Primate, Monkey, Rabbit
Reactivity Notes	Predicted cross-reactivity based on sequence identity: Gorilla (100%), Gibbon (100%), Marmoset (100%), Chicken (88%), Xenopus (82%).
Specificity/Sensitivity	Human DUSP22. BLAST analysis of the peptide immunogen showed no homology with other human proteins, except DUSP15 (59%).
Immunogen	Synthetic 17 aa peptide from near N-terminus of human DUSP22.
Product Application Details	
Applications	Immunohistochemistry, Immunohistochemistry-Paraffin
Recommended Dilutions	Immunohistochemistry, Immunohistochemistry-Paraffin 2 - 22 ug/ml

#### Images

Immunohistochemistry-Paraffin: MKS1 Antibody [NLS6809] - Analysis of anti-DUSP22 antibody with human kidney.





#### **Procedures**

#### Immunohistochemistry Protocol for MKS1 Antibody (NLS6809)

Immunohistochemistry Protocol for MKS1 Antibody (NLS6809): Immunohistochemistry

- 1. Prepare tissue with formalin fixation and by embedding it in paraffin wax.
- 2. Make 4 um sections and place on pre-cleaned and charged microscope slides.
- 3. Heat in a tissue-drying oven for 45 minutes @ 60 degrees Celcius.
- 4. Deparaffinize the tissues by wash drying the slides in 3 changes of xylene for 5 minutes each @ RT.
- 5. Rehydrate the tissues by washing the slides in 3 changes of 100% alcohol for 3 minutes each @ RT.
- 6. Wash the slides in 2 changes of 95% alcohol for 3 minutes each @ RT.
- 7. Wash the slides in 1 change of 80% alcohol for 3 minutes @ RT.
- 8. Rinse the slides in gentle running distilled water for 5 minutes @ RT.

9. Perform antigen retrieval by steaming the slides in 0.01M sodium citrate buffer (pH 6.0) @ 99-100 degrees Celcius for 20 minutes.

- 10. Remove the slides from the heat and let stand in buffer @ RT for 20 minutes.
- 11. Rinse the slides in 1X TBS-T for 1 minute @ RT.

\*\*Do not allow the tissues to dry at any time during the staining procedure\*\*

- 12. Begin the immunostaining by applying a universal protein block for 20 minutes @ RT.
- 13. Drain protein block from the slides and apply the diluted primary antibody for 45 minutes @ RT.
- 14. Rinse the slide in 1X TBS-T for 1 minute @ RT.
- 15. Apply a biotinylated anti-rabbit IgG (H+L) secondary for 30 minutes @ RT.
- 16. Rinse the slide in 1X TBS-T for 1 minute @ RT.
- 17. Apply an alkaline phosphatase steptavidin for 30 minutes @ RT.
- 18. Rinse the slide in 1X TBS-T for 1 minute @ RT.
- 19. Apply an alkaline phosphatase chromagen substrate for 30 minutes @ RT.
- 20. Rinse the slide in distilled water for 1 minute @ RT.

\*\*This method should only be used if the chromagen substrate is alcohol insoluble (ie: Vector Red, DAB)\*\*

21. Dehydrate the tissue by washing the slides in 2 changes of 80% alcohol for 1 minute each @ RT.

- 22. Wash the slides in 2 changes of 95% alcohol for 1 minute each @ RT.
- 23. Wash the slides in 3 changes of 100% alcohol for 1 minute each @ RT.
- 24. Wash the slides in 3 changes of xylene for 1 minute each @ RT.

25. Apply cover slip.





## Novus Biologicals USA

10730 E. Briarwood Avenue Centennial, CO 80112 USA Phone: 303.730.1950 Toll Free: 1.888.506.6887 Fax: 303.730.1966 nb-customerservice@bio-techne.com

## **Bio-Techne Canada**

21 Canmotor Ave Toronto, ON M8Z 4E6 Canada Phone: 905.827.6400 Toll Free: 855.668.8722 Fax: 905.827.6402 canada.inquires@bio-techne.com

## **Bio-Techne Ltd**

19 Barton Lane Abingdon Science Park Abingdon, OX14 3NB, United Kingdom Phone: (44) (0) 1235 529449 Free Phone: 0800 37 34 15 Fax: (44) (0) 1235 533420 info.EMEA@bio-techne.com

#### **General Contact Information**

www.novusbio.com Technical Support: nb-technical@biotechne.com Orders: nb-customerservice@bio-techne.com General: novus@novusbio.com

#### **Products Related to NLS6809**

HAF008	Goat anti-Rabbit IgG Secondary Antibody [HRP]
NB7160	Goat anti-Rabbit IgG (H+L) Secondary Antibody [HRP]
NBP2-24891	Rabbit IgG Isotype Control
NBP2-51697-0.1mg	Recombinant Human DUSP22 His Protein

#### 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.

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