# **Product Datasheet**

# P2Y13/P2RY13/GPR86 Antibody NLS4853

Unit Size: 0.05 ml

Store at 4C short term. Aliquot and store at -20C long term. Avoid freeze-thaw cycles.

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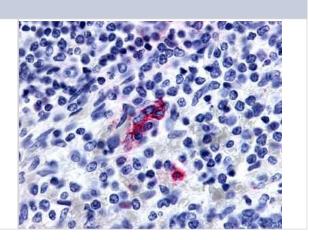
# **NLS4853**

P2Y13/P2RY13/GPR86 Antibody

P2Y13/P2RY13/GPR86 Antibody	
0.05 ml	
1.0 mg/ml	
Store at 4C short term. Aliquot and store at -20C long term. Avoid freeze-thaw cycles.	
Polyclonal	
0.1% Sodium Azide	
IgG	
Immunogen affinity purified	
PBS	
Rabbit	
53829	
P2RY13	
Human	
Predicted cross-reactivity based on sequence identity: Gorilla (100%), Gibbon (93%), Marmoset (93%), Canine (87%), Hamster (80%), Bat (80%), Rabbit (80%), Equine (80%).	
Human P2RY13 / P2Y13. BLAST analysis of the peptide immunogen showed no homology with other human proteins.	
Synthetic 15 amino acid peptide from N-terminal extracellular domain of human P2Y13/P2RY13/GPR86.	
Immunohistochemistry, Immunohistochemistry-Paraffin	
Immunohistochemistry, Immunohistochemistry-Paraffin 11-22 ug/ml	

# **Images**

Immunohistochemistry-Paraffin: P2Y13/P2RY13/GPR86 Antibody [NLS4853] - Analysis of anti-P2RY13 / P2Y13 antibody with human spleen.



### **Procedures**

## Immunohistochemistry Protocol for GPR86 Antibody (NLS4853)

Immunohistochemistry Protocol for GPR86 Antibody (NLS4853): 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 at 60 degrees Celcius.
- 4. Deparaffinize the tissues by wash drying the slides in 3 changes of xylene approximately 5 minutes each @ RT.
- 5. Rehydrate the tissues by washing the slides in 3 changes of 100% alcohol approximately 3 minutes each @ RT.
- Wash the slides in 2 changes of 95% alcohol approximately 3 minutes each @ RT.
- 7. Wash the slides in 1 change of 80% alcohol approximately 3 minutes @ RT.
- 8. Rinse the slides in gentle running distilled water approximately 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 approximately 20 minutes @ RT.
- 13. Drain protein block from the slides and apply the diluted primary antibody approximately 45 minutes @ RT.
- 14. Rinse the slide in 1X TBS-T approximately 1 minute @ RT.
- 15. Apply a biotinylated anti-rabbit IgG (H+L) secondary approximately 30 minutes @ RT.
- 16. Rinse the slide in 1X TBS-T approximately 1 minute at RT.
- 17. Apply an alkaline phosphatase steptavidin approximately 30 minutes at RT.
- 18. Rinse the slide in 1X TBS-T approximately 1 minute at RT.
- 19. Apply an alkaline phosphatase chromagen substrate approximately 30 minutes at RT.
- 20. Rinse the slide in distilled water approximately 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 approximately 1 minute each @ RT.
- 22. Wash the slides in 2 changes of 95% alcohol approximately 1 minute each @ RT.
- 23. Wash the slides in 3 changes of 100% alcohol approximately 1 minute each @ RT.
- 24. Wash the slides in 3 changes of xyleneapproximately 1 minute each @ RT.
- 25. Apply cover slip.





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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

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