Product Datasheet

FPRL1/FPR2 Antibody
NLS1878SS

Unit Size: 0.025 ml

Store at 4C short term. Aliquot and store at -80C long term. Avoid freeze-thaw cycles.

Reviews: 1 Publications: 3

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NLS1878SS

FPRL1/FPR2 Antibody

Product Information	
Unit Size	0.025 ml
Concentration	1.1 mg/ml
Storage	Store at 4C short term. Aliquot and store at -80C long term. Avoid freeze-thaw cycles.
Clonality	Polyclonal
Preservative	0.01% Sodium Azide
Purity	Immunogen affinity purified
Buffer	PBS, 30% glycerol
Target Molecular Weight	38 kDa

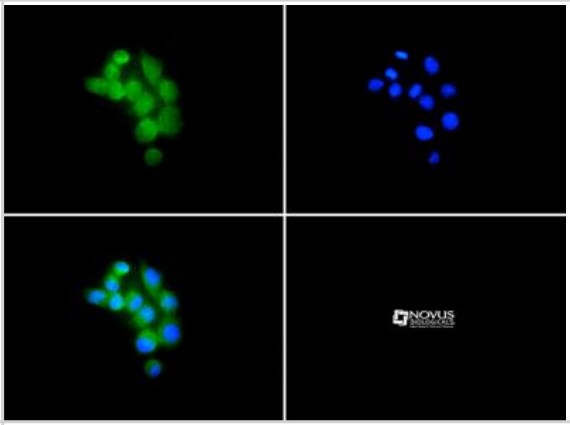
Product Description	
Host	Rabbit
Gene ID	2358
Gene Symbol	FPR2
Species	Human, Mouse, Rat
Species Reactivity	Human and mouse. Rat reactivity reported in scientific literature (PMID: 24086560)
Immunogen	Synthetic peptide made to the 2nd extracellular loop of the human FPRL1 protein (between residues 300-350) [UniProt P25090]

Product Application Details	
Applications	Western Blot, Immunocytochemistry/Immunofluorescence, Immunohistochemistry, Immunohistochemistry-Frozen, Immunohistochemistry-Paraffin
Recommended Dilutions	Immunocytochemistry/Immunofluorescence 1:20-1:75, Immunohistochemistry 1:300, Immunohistochemistry-Frozen, Immunohistochemistry-Paraffin 1:300, Western Blot 1:1000
Application Notes	This FPRL1 antibody is useful for Western Blot, Immunocytochemistry/Immunofluorescence, and Immunohistochemistry paraffin embedded sections. In Western Blot, a band is seen ~38 kDa representing FPRL1. In ICC/IF, plasma membrane staining was observed in Raw264.7 cells. In IHC-P, staining was also observed in the plasma membrane of human kidney cancer tissue. Prior to immunostaining paraffin tissues, antigen retrieval with sodium citrate buffer (pH 6.0) is recommended. Customers have reported success in IF on FFPE mouse kidney tissue, following microwave antigen retrieval. Use in Immunohistochemistry-Frozen reported in scientific literature (PMID 24086560)

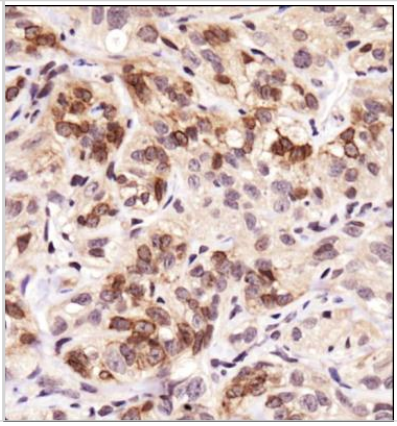
Images

Western Blot: FPRL1 Antibody [NLS1878] - WB analysis of FPRL1 in HL-60 cell lysate.

Immunocytochemistry/Immunofluorescence: FPRL1 Antibody [NLS1878] - FPRL1 antibody was tested in Raw264.7 cells with Dylight 488 (green). Nuclei were counterstained with DAPI (blue).



Immunohistochemistry: FPRL1 Antibody [NLS1878] - IHC analysis of FPRL1 in human kidney cancer using DAB with hematoxylin counterstain.



Publications

Abdelmoaty S, Wigerblad G, Bas DB et al. Spinal Actions of Lipoxin A4 and 17(R)-Resolvin D1 Attenuate Inflammation-Induced Mechanical Hypersensitivity and Spinal TNF Release. PLoS One. 2013 Sep 24 [PMID: 24086560] (IHC-Fr, ICC/IF, Rat)

Chen K, Liu M, Liu Y et al. Formylpeptide receptor-2 contributes to colonic epithelial homeostasis, inflammation, and tumorigenesis J Clin Invest 2013 Mar 1 [PMID: 23454745] (ICC/IF, Mouse)

Singh D, Qi R, Jordan JL et al. The Human Antimicrobial Peptide LL-37, but Not the Mouse Ortholog, mCRAMP, Can Stimulate Signaling by Poly(I:C) through a FPRL1-dependent Pathway. J Biol Chem 2013 Mar 22 [PMID: 23386607] (WB, Human, Mouse)

Procedures

Immunohistochemistry-Paraffin protocol for FPRL1 Antibody (NLS1878)
Immunohistochemistry-Paraffin Embedded Sections

Antigen Unmasking:
Bring slides to a boil in 10 mM sodium citrate buffer (pH 6.0) then maintain at a sub-boiling temperature for 10 minutes. Cool slides on bench-top for 30 minutes.

Staining:
1. Wash sections in deionized water three times for 5 minutes each.
2. Wash sections in wash buffer for 5 minutes.
3. Block each section with 100-400 ul blocking solution for 1 hour at room temperature.
4. Remove blocking solution and add 100-400 ul diluted primary antibody. Incubate overnight at 4 C.
5. Remove antibody solution and wash sections in wash buffer three times for 5 minutes each.
6. Add 100-400 ul biotinylated diluted secondary antibody. Incubate 30 minutes at room temperature.
7. Remove secondary antibody solution and wash sections three times with wash buffer for 5 minutes each.
8. Add 100-400 ul Streptavidin-HRP reagent to each section and incubate for 30 minutes at room temperature.
9. Wash sections three times in wash buffer for 5 minutes each.
10. Add 100-400 ul DAB substrate to each section and monitor staining closely.
11. As soon as the sections develop, immerse slides in deionized water.
12. Counterstain sections in hematoxylin.
13. Wash sections in deionized water two times for 5 minutes each.
14. Dehydrate sections.
15. Mount coverslips.

*The above information is only intended as a guide. The researcher should determine what protocol best meets their needs. Please follow safe laboratory procedures.

Western Blot protocol for FPRL1 Antibody (NLS1878)
Western Blot Protocol

1. Perform SDS-PAGE on samples to be analyzed, loading 40 ug of total protein per lane.
2. Transfer proteins to membrane according to the instructions provided by the manufacturer of the membrane and transfer apparatus.
3. Stain according to standard Ponceau S procedure (or similar product) to assess transfer success, and mark molecular weight standards where appropriate.
4. Rinse the blot.
5. Block the membrane using standard blocking buffer for at least 1 hour.
6. Wash the membrane in wash buffer three times for 10 minutes each.
7. Dilute primary antibody in blocking buffer and incubate 1 hour at room temperature.
8. Wash the membrane in wash buffer three times for 10 minutes each.
9. Apply the diluted 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 three 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.
Note: Tween-20 can be added to the blocking or antibody dilution buffer at a final concentration of 0.05-0.2%.

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Immunocytochemistry/Immunofluorescence Protocol for FPRL1 Antibody (NLS1878)
Immunocytochemistry Protocol

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 1:25,0000 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.

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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 guarantee, please visit www.novusbio.com/guarantee.