# **Product Datasheet**

# CD74 Antibody (PIN.1) - BSA Free NB100-1985SS

Unit Size: 0.025 ml

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

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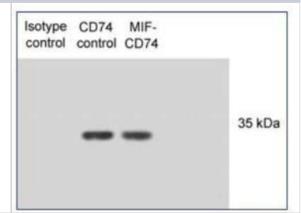
## NB100-1985SS

CD74 Antibody (PIN.1) - BSA Free	
Product Information	
Unit Size	0.025 ml
Concentration	1.0 mg/ml
Storage	Store at 4C short term. Aliquot and store at -20C long term. Avoid freeze-thaw cycles.
Clonality	Monoclonal
Clone	PIN.1
Preservative	0.02% Sodium Azide
Isotype	IgG1
Purity	Protein G purified
Buffer	PBS
Product Description	
Description	Novus Biologicals Mouse CD74 Antibody (PIN.1) - BSA Free (NB100-1985) is a monoclonal antibody validated for use in IHC, WB, Flow, ICC/IF, Simple Western and IP. Anti-CD74 Antibody: Cited in 6 publications. All Novus Biologicals antibodies are covered by our 100% guarantee.
Host	Mouse
Gene ID	972
Gene Symbol	CD74
Species	Human, Mouse (Negative)
Reactivity Notes	Human. Does not react with mouse.
Specificity/Sensitivity	This detects an ~33-35 kDa protein doublet, corresponding to the apparent molecular mass of the p33 and p35 forms of human CD74.
Immunogen	Peptide corresponding to residues 12-27 of CD74, human invariant chain (short form).
Product Application Details	
Applications	Western Blot, Simple Western, Immunohistochemistry-Paraffin, Flow Cytometry, Immunocytochemistry/ Immunofluorescence, Immunohistochemistry, Immunoprecipitation, CyTOF-ready
Recommended Dilutions	Western Blot 1 ug/ml, Simple Western, Flow Cytometry 1 ug/million cells, Immunohistochemistry 5 ug/ml, Immunocytochemistry/ Immunofluorescence 1:10 -1:2000, Immunoprecipitation 1:10-1:500, Immunohistochemistry-Paraffin 5 ug/ml, CyTOF-ready
Application Notes	In Simple Western only 10-15 ul of the recommended dilution is used per data point. In WB this detects an ~33-35 kDa protein doublet, corresponding to the apparent molecular mass of the p33 and p35 forms of human CD74. This antibody is CyTOF ready. See <a href="Simple Western Antibody Database">Simple Western Antibody Database</a> for Simple Western validation: tested in Jurkat lysate (0.2 mg/ml); antibody dilution of 1:400; separated by size; detects a band at 62 kDa

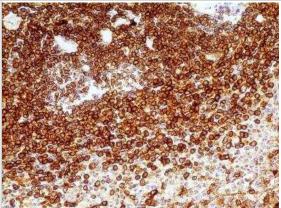


## **Images**

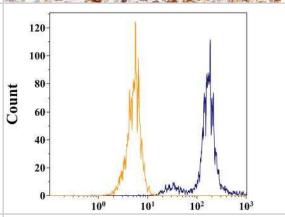
Western Blot: CD74 Antibody (PIN.1) [NB100-1985] - Western blot of CD74 from IP N87 lysates mixed with macrophage migration inhibitory factor (MIF).



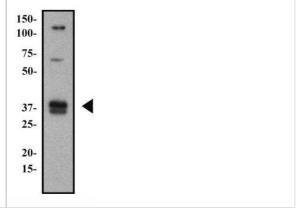
Immunohistochemistry-Paraffin: CD74 Antibody (PIN.1) [NB100-1985] - IHC analysis of formalin-fixed paraffin-embedded tissue section of human spleen using mouse monoclonal CD74 antibody (clone PIN.1) at 5 ug/ml concentration.



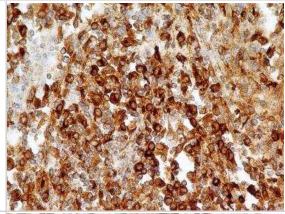
Flow Cytometry: CD74 Antibody (PIN.1) [NB100-1985] - CD74 in human PBMCs (gated on all monocytes). Orange represents isotype control and blue represents anti-CD74 antibody staining.



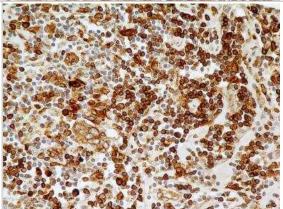
Western Blot: CD74 Antibody (PIN.1) [NB100-1985] - Ramos whole cell protein was separated on a 4-12% gel by SDS-PAGE and protein transferred to PVDF. The membrane was probed with anti-CD74 at 2 ug/ml and detected with and anti-mouse HRP secondary antibody using chemiluminescence. The arrow head shows the detection of CD74.



Immunohistochemistry-Paraffin: CD74 Antibody (PIN.1) [NB100-1985] - IHC analysis of formalin-fixed paraffin-embedded tissue section of human renal cell carcinoma using mouse monoclonal CD74 antibody (clone PIN.1) at 5 ug/ml concentration.



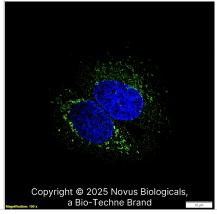
Immunohistochemistry-Paraffin: CD74 Antibody (PIN.1) [NB100-1985] - IHC analysis of formalin-fixed paraffin-embedded tissue section of human pulmonary squamous cell carcinoma using mouse monoclonal CD74 antibody (clone PIN.1) at 5 ug/ml concentration.



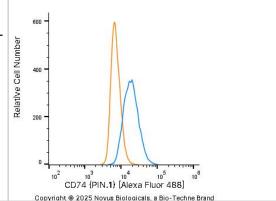
CD74 (PIN.1) was detected in immersion fixed U-2 OS human osteosarcoma cell line using Mouse anti-CD74 (PIN.1) Protein G Purified Monoclonal Antibody conjugated to Alexa Fluor® 647 (Catalog # NB100-1985AF647) (light blue) at 10  $\mu$ g/mL overnight at 4C. Cells were counterstained with DAPI (dark blue). Cells were imaged using a 100X objective and digitally deconvolved.



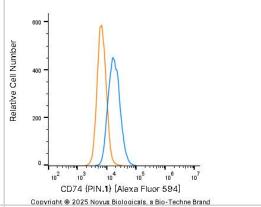
CD74 (PIN.1) was detected in immersion fixed U-2 OS human osteosarcoma cell line using Mouse anti- CD74 (PIN.1) Protein-G purified Monoclonal Antibody conjugated to Alexa Fluor® 488 (Catalog # NB100-1985AF488) (green) at 5 µg/mL overnight at 4C. Cells were counterstained with DAPI (blue). Cells were imaged using a 100X objective and digitally deconvolved.



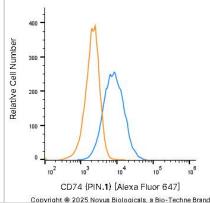
An intracellular stain was performed on THP-1 human acute monocytic leukemia cell line with Mouse anti-CD74 (PIN.1) Protein-G purified Monoclonal Antibody conjugated to Alexa Fluor® 488 (Catalog # NB100-1985AF488, blue histogram) or matched control antibody (orange histogram) at 5 µg/mL for 30 minutes at RT.



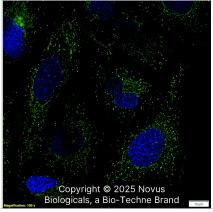
An intracellular stain was performed on THP-1 human acute monocytic leukemia cell line with Mouse anti-CD74 (PIN.1) Protein-G purified Monoclonal Antibody conjugated to Alexa Fluor® 594 (Catalog # NB100-1985AF594, blue histogram) or matched control antibody (orange histogram) at 2.5 µg/mL for 30 minutes at RT.



An intracellular stain was performed on THP-1 human acute monocytic leukemia cell line with Mouse anti-CD74 (PIN.1) Protein-G purified Monoclonal Antibody conjugated to Alexa Fluor® 647 (Catalog # NB100-1985AF647, blue histogram) or matched control antibody (orange histogram) at 5 μg/mL for 30 minutes at RT.



CD74 (PIN.1) was detected in immersion fixed U-2 OS human osteosarcoma cell line using Mouse anti-CD74 (PIN.1) Protein-G purified Monoclonal Antibody conjugated to FITC (Catalog # NB100-1985F) (green) at 2 µg/mL overnight at 4C. Cells were counterstained with DAPI (blue). Cells were imaged using a 100X objective and digitally deconvolved.



### **Publications**

Koh YW, Han JH, Haam S, Lee HW. An immune-related gene expression signature predicts brain metastasis in lung adenocarcinoma patients after surgery: gene expression profile and immunohistochemical analyses Translational Lung Cancer Research 2021-03-15 [PMID: 33718023]

Weng Y, Lou J, Bao Y et al. Single-Cell RNA Sequencing Technology Revealed the Pivotal Role of Fibroblast Heterogeneity in Angiotensin II-Induced Abdominal Aortic Aneurysms DNA and cell biology 2022-04-22 [PMID: 35451888]

Tanese K, Hashimoto Y, Berkova Z et al. Cell Surface CD74-MIF Interactions Drive Melanoma Survival in Response to Interferon-Gamma J. Invest. Dermatol. 2015-06-03 [PMID: 26039541] (WB, Human)

Lamb CA, Cresswell P. Assembly and transport properties of invariant chain trimers and HLA-DR-invariant chain complexes. J Immunol. 1992-06-01 [PMID: 1588042]

Roche PA, Marks MS, Cresswell P. Formation of a nine-subunit complex by HLA class II glycoproteins and the invariant chain. Nature. 1991-12-05 [PMID: 1956401]

Katsel P, Tan W, Haroutunian V. Gain in brain immunity in the oldest-old differentiates cognitively normal from demented individuals. PLoS One 4(10):e7642. 2009-10-29 [PMID: 19865478]



#### **Procedures**

## Immunohistochemistry-Paraffin protocol for CD74 Antibody (NB100-1985)

CD74 Antibody (PIN.1):

- 1. Deparaffinize the tissue sections by immersing the slides in Xylene with two changes for 10 min each. Sections should not get dried at any stage from this point.
- 2. Rehydrate the tissue sections by immersing the slides in decreasing grades of ethanol as follows:
- a. Immerse in 100% ethanol with 2 changes for 5 minutes each
- b. Immerse in 95% ethanol with 2 changes for 5 minutes each
- c. Immerse in 90% ethanol for 5 minutes
- d. Immerse in 70% ethanol for 5 minutes
- e. Immerse in 50% ethanol for 5 minutes
- f. Immerse in distilled water for 5 minutes
- 3. Antigen Retrieval (Microwave Method):
- a. Immerse the slides in a microwave compatible tray containing 10 mM Sodium Citrate buffer (pH 6.0) with 0.05% Tween 20.
- b. Boil the slides and maintain the sub-boiling temperature for 5 minutes in the microwave. Thereafter, take out the tray very carefully and cool it at room temperature (RT) for about 30 minutes.
- c. Wash the slides 3 times, 3 minutes each by immersing them in TBST (Tris Buffered Saline having 0.05% Tween 20).
- 4. Quenching of Endogenous Peroxidase:
- a. Incubate the slides in 3% hydrogen peroxide prepared in methanol for 15 minutes (at RT, in dark conditions).
- b. Wash the slides in TBST 3 times, 3 minutes each.
- 5. Protein Blocking:
- a. Incubate the sections with background sniper solution at RT for 15 minutes (Biocare Medicals, USA).
- b. Wash the sections 3 times, 3 min each by immersing the slides in TBST.
- 6. Primary Antibody:
- a. Dilute the primary antibody at 5ug/ml concentration using PBS as a diluent.
- b. Incubate the sections with diluted primary antibody for 90 minutes at RT in a humidified chamber.
- c. Thereafter, wash the slides 4 times, 5 minutes each with TBST.
- 7. Probe (Secondary Reagent):
- a. Incubate with MACH 1 Mouse probe for 15 minutes at RT.
- b. Incubate for 30 min at room temperature with HRP-Polymer (Biocare Medical, USA).
- c. Wash the slides with TBST 4 times, 5 minutes each
- 8. Chromogen:
- a. Mix 32ul of DAB Chromogen with 1 ml of DAB substrate buffer (Biocare Medical, USA).
- a. Apply 200ul DAB mixture/section and incubate at RT in dark conditions (few seconds 5 minutes).
- b. As soon as an appropriate color develops, rinse the slides with deionized water (2-3 brief rinses).
- 9. Counter stain:
- a. Counter stain with Hematoxylin for 30 seconds (Vector Labs, USA).
- b. Wash in deionized water for 1-2 minutes to clear the extra stain.
- c. Incubate the slides in bluing solution or Scott's water twice for 2 minutes each time.
- 10. Dehydrate the sections in increasing grades of alcohols:
- a. 50% alcohol for 1 minute
- b. 70% for 1 minute
- c. 90% for 1 minute
- d. 95% for 1 minute
- e. 100% for 1 minute
- f. Xylene with 2 changes for 2 minutes each
- 11. Mount with DPX mount and cover-slip glass (Fisher Scientific, USA), carefully not allowing any air bubbles to enter.

NOTE:- This protocol is provided as a reference tool only. Depending upon the type of tissues /tissue processing and reagents employed, the end user will need to optimize the final conditions for achieving an expected staining.





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### Limitations

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