



PolyStain DS Kit - For Mouse and Rabbit antibody on Mouse tissue

(BCIP/AEC)

NB-23-00095- 3(120 ml)

NB-23-00095- 2(36 ml)

NB-23-00095- 1(12 ml)



PolyStain DS Kit - for Mouse and Rabbit antibody on Mouse tissue (BCIP/AEC)

NB-23-00095-1; NB-23-00095-2; NB-23-00095-3

Storage: 2-8°C

INTENDED USE:

The PolyStain DS Kit is designed to use with user supplied mouse and rabbit primary antibody to detect two distinct antigens on mouse tissue or cell samples. NB-23-00090 kits can be used on frozen or paraffin embedded tissues, and freshly prepared monolayer cell smears. Our system is designed not give background on most mouse strains however there may be some mouse strains especially when using frozen that require additional blocking; we recommend PureStain Mouse-on-Mouse Kit Blocking A & B solutions (NB-23-00076) to improve specificity of the mouse primary antibody on mouse tissue. Double staining is one of most common methods used in immunohistostaining that allows for revealing two distinct antigens in a single tissue. The PolyStain DS Kit from Neobiotech Labs supplies two polymer enzyme conjugates: Mouse AP Polymer and Rabbit HRP Polymer with two distinct substrates/chromogens, BCIP/NBT (purple color, use with the Mouse AP Polymer) and AEC (red color, use with the Rabbit HRP Polymer). A Primer step is used to increase specificity of antibody staining. Both enzyme conjugates are applied to the specimen at the same time and mixed on the slide. This kit offers simplified steps that make for a quicker and easier protocol than that used in a sequential procedure. The PolyStain DS Kit is non-biotin system that avoids endogenous biotin non-specific binding.

KIT COMPONENTS:

Component No.	Content	12mL Kit	36mL Kit	120mL Kit
Reagent 1	Mouse Primer (RTU)	12ml	18mlx2	120ml
Reagent 2	Mouse AP Polymer (RTU)	6ml	18ml	60ml
Reagent 3	Rabbit HRP Polymer (RTU)	6ml	18ml	60ml
Reagent 4	BCIP/NBT Chromogen(RTU)	15ml	18mlx2	120ml
Reagent 5A	AEC Substrate (20x)	1ml	2ml	6ml
Reagent 5B	AEC Chromogen (20x)	2ml	4ml	12ml
Reagent 5C	Hydrogen Peroxide (20x)	1ml	2ml	6ml
Reagent 6	NeoMount Universal (RTU)	15ml	18mlx2	120ml

RECOMMENDED PROTOCOL:

1. Fixation: To ensure the quality of the staining and obtain reproducible performance, user needs to supply appropriately fixed tissue and well prepared slides.
2. Tissue needs to be adhered to the slide tightly to avoid falling off.
3. Paraffin embedded sections must be deparaffinized with xylene and rehydrated with a graded series of ethanol before staining.
4. Cell smear samples should be made up to as much of a monolayer as possible to obtain satisfactory results.
5. Three control slides will aid the interpretation of the result: positive tissue control, reagent control (slides treated with Isotype control reagent), and negative control.
6. Proceed IHC staining: **DO NOT** let specimen or tissue dry from this point on.
7. We recommend TBS-T to be used as the wash buffer to get the highest sensitivity and clean background. Phosphate in the PBS-T may inhibit the activity of the alkaline phosphatase. Note: 1X TBS-T =50mM Tris HCl, 150mM NaCl, 0.05% Tween-20 pH7.6.

Reagent	Staining Procedure	Incubation Time (Min.)
1. Peroxidase and Alkaline Phosphatase Blocking Reagent Not provided Fast, easy and it will block endogenous alkaline phosphatase	a. Incubate slides in peroxidase and alkaline phosphatase blocking reagent. We recommend NeoPure Dual Enzyme Block NB-23-00193 . b. Rinse the slide using distilled water.	10 min.
2. HIER Pretreatment: Refer to Ab data sheet.	a. Heat Induced Epitope Retrieval (HIER) may be required for primary antibody suggested by vendor. b. Wash with PBS-T containing 0.05% Tween-20 or 1X TBS-T (See note 7 above) ; 3 times for 2 minutes each.	
3. PureStain Mouse-on-Mouse Kit Blocking A & B solutions (NB-23-00076) (optional see protocol note 2)	a. Add 2 drops (100µl) or enough volume of PureStain Mouse-on-Mouse Kit Blocking A & B solutions to cover the tissue section and Incubate. b. Rinse with PBS with 0.05% Tween-20 for 2 min., 3 times	30 min.

<p>4. PureStain Mouse-on-Mouse Kit Blocking A & B solutions (NB-23-00076) Not provided (optional see protocol note 2)</p>	<p>a. Add 2 drops (100µl) or enough volume of PureStain Mouse-on-Mouse Kit Blocking A & B solutions to cover the tissue section and Incubate. Do not exceed 5min. b. Rinse with PBS with 0.05% Tween-20 for 2 min., 3 times.</p>	<p>5 min.</p>
<p>5. Mouse antibody 1 and Rabbit antibody 2: Supplied by user</p>	<p>Note: Investigator needs to optimize titer of primary antibody and incubation time prior to double staining. a. Apply 2 drops or enough volume of both Mouse primary Antibody 1 and Rabbit primary Antibody 2 to cover the tissue completely. Mix well on the slide and Incubate in moist chamber for 30-60 min. b. Wash with PBS-T containing 0.05% Tween-20 or 1X TBS-T; 3 times for 2 minutes each.</p>	<p>30-60 min</p>
<p>6. Reagent 1 Mouse Primer (RTU)</p>	<p>a. Add 1-2 drops (100µl) or enough volume of Reagent 1 (Mouse Primer) to cover the tissue section and Incubate Room Temperature for 10-15minutes. b. Wash with PBS-T containing 0.05% Tween-20 or 1X TBS-T; 3 times for 2 minutes each.</p>	<p>10-15min</p>
<p>7. Mix Reagent 2&3 Reagent 2 : Mouse AP Polymer (RTU) Reagent 3: Rabbit HRP Polymer (RTU)</p>	<p>Note: Make sufficient polymer mixture by adding Reagent 2 (Mouse AP Polymer) and Reagent 3 (Rabbit HRP Polymer) at 1:1 ratio, mix well. Do Not mix more than you need for the experiment because the polymer mixture is not stable for long term storage. a. Apply 1 to 2 drops (50-100µl) of the mixture to cover the tissue completely. b. Incubate in moist chamber for 30 min. c. Wash with 1X TBS-T only; 3 times for 2 minutes each.</p>	<p>30 min</p>
<p>8. Reagent 4 BCIP/NBT Chromogen (RTU)</p>	<p>a. Apply 2 drops or enough volume of Reagent 4 (BCIP/NBT Chromogen) to completely cover tissue. Incubate for 3-10 min. b. Rinse thoroughly with distilled water. c. Wash with PBS-T containing 0.05% Tween-20 or 1X TBS-T; 3 times for 2 minutes each.</p>	<p>3-10 min</p>
<p>9. Reagent 5A, 5B, 5C: 5A:AEC Substrate Buffer (20x) 5B:AEC Chromogen (20x) 5C:Hydrogen Peroxide (20x)</p>	<p>a. Add 1 drop (50µl) of Reagent 5A to 1ml distill water. Mix well. Add 2 drops of Reagent 5B and 1 drop of Reagent 5C to diluted AEC Substrate. Mix well. Keep away from light and use within 1 hour. b. Apply 2 drops (100µl) or enough volume of AEC working solution to completely cover the tissue. Incubate for 5-15 min, observe appropriate color development. c. Rinse well with distilled water. (AEC is alcohol soluble; do not dehydrate.)</p>	<p>5-15 min.</p>

<p>10. HEMATOXYLIN Not provided</p>	<p>a. Counterstain with 2 drops (100µl) or enough volume of hematoxylin to completely cover tissue. Incubate for 10-15 seconds.</p> <p>b. Rinse thoroughly with tap water for 2-3 min.</p> <p>c. Put slides in PBS until show blue color (about 30 - 60 sec.)</p> <p>d. Rinse well in distilled water</p>	
<p>11. Reagent 6: NeoMount Universal (RTU)</p> <p>To coverslip see protocol note 3</p>	<p>a. Apply 2 drops (100µl) or enough volume of Reagent 6 (NeoMount Universal) to cover tissue when tissue is wet. Rotate the slides to allow NeoMount Universal spread evenly. DO NOT coverslip.</p> <p>b. Place slides horizontally in an oven at 40-50°C for at least 30 minutes or leave it at room temperature until slides are thoroughly dried. Hardened NeoMount Universal forms an impervious polymer barrier to organic solvent. Do not use oil directly on the top of dried NeoMount Universal</p>	<p>30 min. in 40-50°C oven Or: overnight at room temperature</p>

PROTOCOL NOTES:

1. The fixation, tissue slide thickness, antigen retrieval and primary antibody dilution and incubation time affect results significantly. Investigator needs to consider all factors and determine optimal conditions when interpreting the result.
2. Mouse-on-Mouse Kit Blocking (sample provided) the anti-mouse secondary has been absorbed to rat serum resulting in most mouse strains having no background, however some mouse strains may need additional blocking. Mouse-on-Mouse Kit Blocking (NB-23-00076) works very well on frozen tissue.
3. NeoMount Universal is an aqueous-based mounting media for immunohistochemistry. It is used as the permanent mounting media for alcohol soluble chromogens such as AP-Red, AEC, and BCIP. NeoMount Universal does not use a coverslip. However, if you need to coverslip your tissue, after NeoMount Universal has dried, dip the slide in xylene (1 to 2 seconds), apply an organic mounting solution (such as NeoMount Perm, Cat# NB-23-00156), and place cover glass on the slide. Store slides after they have dried completely.

PRECAUTIONS:

Please wear gloves and take other necessary precautions.

FOR RESEARCH USE

Work Sheet for NB-23-00095 Kit

We designed this work sheet to help you keep track of each step. We recommend you use this sheet to record the actual time of each step conducted as it will be helpful for questions with our technical support.

- Used for tester to check “√” each step during the experiment
- Steps follow after de-paraffinization
- Refer to insert for details of each step

Protocol Step	NB-23-00095 Protocol	Experiment 1 Date:	Experiment 2 Date:	Experiment 3 Date:	Experiment 4 Date:
Step 1	Peroxidase Block				
Step 2	HIER if needed				
Step 3 Optional	PureStain Mouse-on-Mouse Kit Blocking A 30 min. NB-23-00076				
Step 4 Optional	PureStain Mouse-on-Mouse Kit Blocking B 5 min. NB-23-00076				
Step 5	Ms 1°Ab & Rb 1°Ab mix (30-60 min.)				
Step 6	Reagent 1 Mouse Primer (15 min.)				

Step 7	Reagent 2 & Reagent 3 Ms AP Polymer & Rb HRP Polymer require mixing (30 min.)				
Step 8	Reagent 4 BCIP/NBT (3-10 min.)				
Step 9	Reagent 5A, 5B, & 5C AEC Requires mixing! (5- 15 min.)				
Step 10	Counter stain Hematoxylin User supplied				
Step 11	Reagent 6 Simpomount Do not coverslip!				

Testing result: