NeoStain ABC Kit, AP Detection Kit for Mouse Antibodies

NB-23-00024

NeoStain ABC Kit, Alkaline Phosphatase Detection Kit for Mouse Antibodies

(Alkaline phosphatase labeled streptavidin-biotin detection system for mouse antibody)

#Cat: NB-23-00024-1	**Size: 110ml, no chromogen**
#Cat: NB-23-00024-2	**Size: 18ml, with Fast Red**
#Cat NB-23-00024-3	**Size: 6ml, with Fast Red**
#Cat: NB-23-00024-4	**Size: 18ml, with Permanent Red**
#Cat: NB-23-00024-5	**Size: 6ml, with Permanent Red**

Intended Use:

NeoStain AP Mouse Detection Kit uses biotinylated secondary antibody and Alkaline Phosphatase (AP) labeled-streptavidin to detect mouse primary antibody (user-supplied) that bind to antigens in human tissue or cell preparations under light microscopy. The most commonly used specimens for this system are: frozen tissue, paraffin-embedded tissue, freshly prepared lymphocytes and fixed culture cells. Alkaline Phosphatase (AP) labeled-streptavidin and biotinylated secondary antibody amplification system has become a standard technique in immunochemical staining1,2. NeoStain AP Mouse Detection Kit uses human-absorbed, bioinylated, affinity-purified secondary antibody reacts with the user supplied primary antibody bound to the specific epitope of the antigen in tissue or cell. Alkaline Phosphatase (AP) labeled streptavidin then reacts with biotinylated secondary antibody to form an AP-streptavidin-biotin complex. The AP enzyme of the streptavidin complex catalyzes the substrate/chomogen such as Fast-Red, AP-Red, or BCIP/NBT to form a red (Fast-Red or AP-Red) or dark blue/purple (BCIP/NBT) color deposit at the antigen site. The antigen then can be visualized under microscope. Compared to traditional ABC methods which uses avidin, NeoStain AP Broad Detection Kit demonstrates stronger binding strength to bind biotin and less non-specific background staining.

Higher sensitivity and lower background give this kit a higher signal-noise ratio. It also provides users cost effective method for their research. End users may choose Fast-Red, AP-Red, or BCIP/NBT chromogen depending on their preferences.

Kit Components:

Component No.	Content	NB-23-00024-5	NB-23-00024-4	NB-23-00024-1
Reagent 1	Pre-Block Solution (RTU)	6mL	18mL	110mL
Reagent 2	Biotinylated anti-Mouse (RTU)	6mL	18mL	110mL
Reagent 3	Streptavidin-AP (RTU)	6mL	18mL	110mL
Reagent 4A	Permanent Red Substrate (RTU)	7mL	18mL	NA
Reagent 4B	Permanent Red Activator (5x)	1.4mL	3.6mL	NA
Reagent 4C	Permanent Red Chromogen (100x)	70µL	180µL	NA

Component No.	Content	NB-23-00024-3	NB-23-00024-2
Reagent 1	Pre-Block Solution (RTU)	6mL	18mL
Reagent 2	Biotinylated anti-Mouse (RTU)	6mL	18mL
Reagent 3	Streptavidin-AP (RTU)	6mL	18mL
Reagent 4A	Fast Red tablets (Tablets)	6 tablets	15 Tablets
Reagent 4B	Fast Red Substrate (RTU)	35mL	80mL

Reagent	Staining Procedures	Incubation Time
1. HIER Pretreatment: refer to antibody spec. sheet	a. Heat Induced Epitope Retrieval (HIER) may be required for primary antibody. Refer to antibody datasheet. b. Wash with PBS-T containing 0.05% Tween-20 or **1X TBS-T** **(See note 7 above)**; 3 times for 2 minutes each.	
2. **Reagent 1:** Pre-blocking Solution (RTU)	a. Add 2 drops or enough volume of **Regent 1** (Pre-blocking Solution) to completely cover the tissue section and incubate for 10 min. b. Blot off solution. **DO NOT RINSE**.	1min.
3. Primary antibody: Supplied by user.	Note: Investigator needs to optimize dilution and incubation time. a. Apply 2 drops or enough volume of Primary antibody to cover the tissue section completely.Incubate in moist chamber for 30-60min. b. Wash with PBS-T containing 0.05% Tween-20 or **1X TBS-T**; 3 times for 2 minutes each.	30-60min.
4. **Reagent 2:** Biotinylated anti-Mouse (RTU)	a. Apply 2 drops or enough volume of **Reagent 2** (Biotinylated anti-Mouse) to cover the tissue section completely and incubate for 10min. b. Wash with PBS-T containing 0.05% Tween-20 or **1X TBS-T**; 3 times for 2 minutes each.	10min.
5. **Reagent 3:** Streptavidin-AP (RTU)	a. Apply 2 drops or enough volume of **Reagent 3** (Streptavidin-AP)to cover the tissue section completely and incubate for 10min. b. Wash with **1xTBS-T** only, 3 times for 2 minutes each.	10min.
6. **Reagent 4:** Chromogen: Fast-Red, or Permanent Red **To get maximum sensitivity of AP polymer, Please repeat chromogenstep**), or BCIP.NBT	Refer to manufacture data sheet if chromogen is supplied by user. Recommended protocol for chromogen using our kit: **1. Fast Red :** a. Dissolve one Fast Red tablet into one 5mL substrate buffer. Vortex until tablet is dissolved. Itusually takes 20 minutes to dissolve completely. b. Chromogen must be used within 1 hour. c. Apply 100ul or more Fast-Red solution to completely cover the tissue section and incubate 10minutes at room temperature. d. After proper color development, wash with distill water for 2 minutes, 3 times e. DO NOT Dehydrate tissue after staining. Fast-Red is alcohol soluble. **2. Permanent Red:** **Note**: Shake Permanent Red Activator before adding into Permanent Red Substrate. a. Add 200µL of Reagent 7B (Activator) into 1mL of Reagent 7A (Substrate) and mix well. Add10µL of Reagent 7C (Chromogen) into the mixture and mix well. [Note: For fewer slides, Add 100µL of Reagent 7B (Activator) into 500µL of Reagent 7A(Substrate) and mix well. Add 5µL of Reagent 7C (Chromogen) into the mixture and mix well.] **b.** Apply 2 drops (100µL) or enough volume of Permanent Red working solution to completely cover the tissue. Incubate for 10 min, observe appropriate color development. **To increase AP signal aspirate or tap off chromogen and apply 2-3 drops (100µL) again of the Permanent Red working solution to completely cover the tissue for additional 5 to 10min.** c. Rinse well with distilled water. **3. BCIP/NBT :** order separately, Cat.# NB-23-00144-1 / -2 a. Add two drops (about 100ul) of Ready-to-use BCIP/NBT to cover the tissue section for 5-10 minutes. Monitor the color development under a microscope. b. Rinse with distill water for 2 minutes, 3 times.	
7. Hematoxylin: Supplied by user	a. Counterstain with 2 drops or enough volume to cover tissue completely and wait about 10-20 seconds. b. Rinse thoroughly under tap water for 1-2 min. c. Put slides in PBS until show blue color (about 30-60 seconds) d. Rinse well in distilled water	
8. Mountinga: Supplied by user	Follow the manufacturer's data sheet procedure for mounting. Recommended product: 1. NeoBio Mount AQ Cat.# NB-23-00155-3 (18ml) for AEC, Fast-red, AP-Red and AP-blue, DAB, BCIP/NBT. 2. NeoBio Mount Perm: Cat.# NB-23-00156 (18ml), for DAB and BCIP/NBT 3. NeoBio Mount Universal: Cat.# NB-23-00157-2 (18ml), or NB-23-00157-1 (100ml), universal permanent mounting medium	

Recommended Protocol:

1. Fixation: To ensure the quality of the staining and obtain reproducible performance, the user needs to supply appropriately fixed tissue and well prepared slides.
2. Tissue need to be adhered to the slide tightly to avoid tissue falling off.
3. Paraffin embedded section must be deparffinized with xylene and rehydrated with a graded series of ethanol before staining.
4. Cell smear samples should be made into as thin monolayer as possible to obtain satisfactory results.
5. Three control slides will aid the interpretation of the result: positive tissue control, reagent control (slide treated with Isotype control reagent), and negative control.
6. Start staining procedures: 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. (10xTBS-T NB-23-00201)

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. Tissue staining is dependent upon the proper handling and processing of tissues prior to staining. Improper tissue preparation may lead to false negative results or inconsistent results.
3. Do not mix reagents from different lot.
4. Do not allow the slides to dry at any time during staining.

Precautions:

Handle all specimens as potential infectious materials, wear gloves and protection cloth when handling all reagents.

Storage: Store at 4°C.

References:

1. Elias, J.M. et al. Sensitivity and Detection Efficiency of the Peroxidase antiperoxidase (PAP) Avidin-Biotin Peroxidase Complex (ABC), and Peroxidase-Labeled Avidin-Biotin (LAB Methods. AM J Clin Pathol 92:62-67, 1989.
2. Polak, J.M and Van Noorden, S. Introduction to Immunocytochemistry Second Edition. Bios Scientific Publishers. 41-54. 1997.

Remarks:

For research use or investigation only. Not for diagnostic or therapeutic use.

Related Products:

NeoStain ABC Kit AP Board Bulk	NB-23-00023-1	110ml
NeoStain ABC Kit AP Board Fast Red	NB-23-00023-2 / -3	18ml / 6ml
NeoStain ABC Kit AP Board AP-Red+	NB-23-00023-4 / -5	18ml / 6ml
NeoStain ABC Kit AP Rabbit Bulk	NB-23-00025-1	110ml
NeoStain ABC Kit AP Rabbit Fast Red	NB-23-00025-2 / -3	18ml / 6ml
NeoStain ABC Kit AP Rabbit AP-Red+	NB-23-00025-4 / -5	18ml / 6ml
Streptavidin-AP (RTU)	NB-23-00027-1 / -2	110ml / 18ml
Fast Red Kit	NB-23-00142	12 Tab + 60ml
AP-Red+ Kit (40x concentrate)	NB-23-00143	8ml
BCIP/NBT Kit	NB-23-00144-1 / -2	100ml / 18ml
NeoBio Mount AQ (Aqueous)	NB-23-00142-3	18ml
NeoBio Mount Perm (Organic)	NB-23-00156	18ml
NeoBio Mount Universal (Aqueous)	NB-23-00157-1 / -2	100ml / 18ml