

<u>NeoStain ABC Kit, HRP</u> <u>Detection Kit for Mouse</u> <u>Antibodies</u>

NB-23-00003



NeoStain ABC Kit for Mouse Antibodies

(Horseradish peroxidase labeled-streptavidin-biotin detection system for mouse antibody with DAB chromogen)

#Cat: NB-23-00003-1	Size: 500mL, no chromogen
#Cat: NB-23-00003-2	Size: 110ml, no chromogen
#Cat: NB-23-00003-3	Size: 18ml, with DAB
#Cat: NB-23-00003-4	Size: 6ml, with DAB

Intended Use:

NeoStain HRP Mouse Detection (DAB) kit is intended for using with mouse primary antibody (usersupplied) to detect the presence of antigens in human tissue or cell preparations under light microscopy. Most commonly used specimens for this system are: frozen tissue, paraffin-embedded tissue, freshly prepared lymphocytes and fixed culture cells.

Horseradish peroxidase (HRP) labeled-streptavidin and biotinylated secondary antibody amplification system has become a standard technique in immunochemical staining1,2. NeoStain HRP Mouse Detection (DAB) kit uses human absorbed, biotinylated, affinity-purified secondary antibody reacts with the user supplied primary antibody bound to the specific epitope of the antigen in tissue or cell. Horseradish peroxidase (HRP) labeled streptavidin then reacts with biotinylated secondary antibody to form a HRP-streptavidin-biotin complex. The HRP enzyme of the streptavidin complex catalyzed the substrate/chomogen, 3,3' diaminobenzidine (DAB substrate) reaction to form brown color deposit at the antigen site. The antigen then can be visualized under microscope. Compared to traditional ABC method which uses avidin, NeoStain HRP Mouse Detection (DAB) kit demonstrates stronger binding strength to bind biotin and less non-specific background staining. Pre-Blocking Solution in the kit will help to eliminate non-specific background.

Higher sensitivity and lower background give NeoStain HRP Mouse Detection (DAB) kit a higher signalnoise ratio. More than sufficient volume of DAB chormogen is provided in the kit so that customers may use 2 drops of DAB chromogen per ml to obtain higher sensitivity and contrast.

		Reagent 1	Reagent 2	Reagent 3	Reagent 4A	Reagent 4B
Cat. #	Name	Pre-Blocking	Biotinylated anti-	Streptavidin	DAB	DAB
		Solution	mouse secondary	peroxidase	substrate	chromogen
			antibody	conjugate	(Ready-to-	(Concentrated)
					use)	
NB-23-00003-4	NeoStain ABC Kit,	6ml	6 ml	6 ml	12 ml	1.5ml
	HRP, Mouse, DAB					
NB-23-00003-3	NeoStain ABC Kit,	18 ml	18 ml	18 ml	15ml x 2	2 ml
	HRP, Mouse, DAB					
NB-23-00003-2	NeoStain ABC Kit,	110ml	110ml	110ml	Not	Not included
	HRP, Mouse, no				included	
	chromogen					
NB-23-00003-1	NeoStain ABC Kit,	500ml	500ml	500ml	Not	Not included
	HRP, Mouse, no				included	
	chromogen					

Kit Components:



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 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 as much 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.

Reagent	Staining Procedures	Incubation
		Time (Min.)
 Peroxidase blocking reagent: Supplied by user. We recommend using Peroxidase Block NB-23-00192- 1 /-2. 	 a. Apply 2 drops (100 μL) or enough volume of Peroxidase blocking reagent (Ready-to-use 3% H2O2 solution) to cover the tissue section and incubate b. Rinse the slide using distilled water. 	10 min.
2. HIER Pretreatment: refer to antibody spec. sheet	a. Heat Induced Epitope Retrieval (HIER) may be required for primary antibody suggested by vendor b. b. Wash with PBS 2 min., 3 times.	
3. Reagent 1: Pre-blocking Solution	a. Add 2 drops or enough of volume Pre-blocking Solutionto completely cover the tissue section and Incubateb. Blot off solution. DO NOT RINSE.	10 min.
 4. Primary antibody: Supplied by user. 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 30-60 min. b. Rinse with PBS for 2 min., 3 times. 	30-60 min.
5. Reagent 2: Ready to use Secondary antibody	a. Apply 2 drops or enough volume of secondary antibodyto cover the tissue section completely and incubate.b. Rinse with PBS for 2 min., 3 times.	10 min.
6. Reagent 3: Ready to use HRP-Streptavidin	a. Apply 2 drops or enough volume of HRP-Streptavidin to cover the tissue section completely and incubate.b. Rinse with PBS for 2 min., 3 times.	10 min.
 7. Reagents 4A, 4B: 4A: DAB Substrate (RTU) 4B: DAB Chromogen Concentrate (chromogen may be supplied by user) 	 a. Add 1 drop or 2 drops (for higher sensitivity and contrast) of Reagent 4B to 1ml of 4A. Mix well. Protect from light and use within 5 hours. b. Apply 2 drops (100 μL) or enough volume of pre-mixed DAB chromogen to completely cover tissue and Incubate 5 minutes. c. Rinse with distill water for 2 min, 3 times. 	5 min.



8. Hematoxylin:	a. Counterstain with 2 drops or enough volume to	
Supplied by user	covertissue 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-	
	60seconds)	
	d. Rinse well in distilled water	
9. Mounting	Follow the manufacture data sheet procedure for	
media:	mounting. Recommended product:	
Supplied by user	1. NeoBio Mount Perm: Cat.# NB-23-00156 for DAB	
	2. NeoBio Mount Universal: Cat.# NB-23-00157-2	
	(18ml), or -1 (100ml), universal permanent mounting	
	medium	

Protocol Notes:

- 1. The fixation, tissue slide thickness, antigen retrieval and primary antibody dilution and incubation time effect results significantly. Investigator needs to consider all factors and determine optimal conditions when interpret 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

Precautious:

DAB may be carcinogenic. Handle all specimens as potential infectious materials, wear gloves and protection cloth.

Remarks:

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

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.



Related Products:

Product	Catalog No.	Size	Product	Catalog No.	Size
NeoStain ABC Kit, HRP, Rabbit & Mouse, no chromogen	NB-23-00001-3	110mL	Simplified HRP Rabbit Kit (Concentrated, suggested 1:100-200)	NB-23-00010	1 mL
NeoStain ABC Kit, HRP, Rabbit & Mouse, with DAB	NB-23-00001-5 NB-23-00003-6	18 mL 6 mL	Simplified HRP Mouse Kit (Concentrated, suggested 1:100-200)	NB-23-00011	1 mL
NeoStain ABC Kit, HRP, Rabbit, no chromogen	NB-23-00005-2	110mL	Streptavidin-HRP (RTU)	NB-23-00026-2 NB-23-00026-3	18 mL 6 mL
NeoStain ABC Kit, HRP, Rabbit, with DAB	NB-23-00005-3 NB-23-00005-4	18mL 6mL	NeoStain ABC Kit, HRP, Mouse & Rabbit, with AEC	NB-23-00007-1 NB-23-00007-2	18 mL 6 mL
NeoStain ABC Kit, HRP, Goat, no chromogen	NB-23-00012-1	110mL	NeoStain ABC Kit, HRP, Mouse, with AEC	NB-23-00008-1 NB-23-00008-2	18 mL 6 mL
NeoStain ABC Kit, HRP, Goat, with DAB	NB-23-00012-2 NB-23-00012-3	18 mL 6 mL	NeoStain ABC Kit, HRP, Rabbit, with AEC	NB-23-00009-1 NB-23-00009-2	18 mL 6 mL