



NeoStain Poly 1-Step Kit,
HRP Detection System for
Goat Antibodies
(for DAB)

NB-23-00031

NeoStain Poly 1-Step Kit, Horseradish peroxidase Detection System Kit for Goat Antibodies (for DAB)

(NeoStain-HRP detection system, biotin-free, Anti-goat)

Ready-to-use One Step Polymer Detection System

#Cat: NB-23-00031-1	Size: 110mL, no chromogen
#Cat: NB-23-00031-2	Size: 18ml, with DAB (good for 150 slides)
#Cat: NB- 23-00031-3	Size: 6ml, with DAB (good for 50 slides)

Intended Use:

NeoStain Poly 1-Step Goat DAB Detection Kit is designed to use with user supplied goat antibody to detect target antigen on human tissue or cell samples. Specimen can be frozen or paraffin–embedded tissues, and freshly prepared monolayer cell smears.

NeoStain Poly 1-Step Goat DAB Detection Kit is the ONE step polymer detection system that uses polymeric horseradish peroxidase (HRP) -linked goat anti goat IgG to directly detect primary antibody that bound to the tissue. This technology provides excellent sensitivity and high specificity. It is a biotin-free system, therefore, overcomes the non-specific staining caused by streptavidin/biotin system due to endogenous biotin¹. It is a ONE step detection system that is much faster assay compared to traditional two step method (Biotinylated 2nd antibody, and then streptavidin-HRP). These advantages provide laboratories the benefit of more accurate and quicker result, less trouble shooting and better cost-saving. For AEC staining please choose NeoStain Poly 1-Step HRP Goat for AEC (NB-23-00036-1).

Kit Components:

Catalog No.	Product Name	Reagent 1: Polymer HRP-linked anti-goat IgG (Ready-to-use)	Reagent 2: 2A: DAB Substrate 2B: Chromogen concentrate
NB-23-00030-1	NeoStain Poly 1-Step no chromogen	110ml	Not provided
NB-23-00030-2	NeoStain Poly 1-Step with DAB	18ml	30 ml of 2A and 2 ml of 2B
NB-23-00030-3	NeoStain Poly 1-Step with DAB	6ml	12 ml of 2A and 1.5 ml of 2B

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 deparaffinized 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. Investigator needs to optimize dilution and incubation times for primary antibodies.
6. Three control slides will aid the interpretation of the result: positive tissue control, reagent control (slides treated with Isotype control reagent), and negative control.
7. Proceed IHC staining: DO NOT let specimen or tissue dry from this point on.

Reagent:

Reagent	Staining Procedure	Incubation Time (Min.)
1. Peroxidase Blocking Reagent Supplied by user	<ul style="list-style-type: none"> a. Incubate slides in peroxidase blocking reagent (Ready-to-use 3% H₂O₂ solution) for 10 min. b. Rinse the slide using distilled water. 	10
2. HIER Pretreatment: Refer to antibody data sheet.	<ul style="list-style-type: none"> a. Heat Induced Epitope Retrieval (HIER) may be required for primary antibody suggested by vendor. b. Wash with PBS 3 times for 2 minutes each time. 	Refer to vendor's data sheet
3. Pre-Block (Optional) Not provided	<ul style="list-style-type: none"> a. Add 2 (100 µL) or more drops of 10% Normal Goat Serum to cover the tissue section and Incubate 10 min. b. Drain or blot off solution. DO NOT RINSE. 	10
4. Primary antibody: Supplied by user	<p>Notes: Investigator needs to optimize dilution and incubation times</p> <ul style="list-style-type: none"> a. Apply 2 (100 µL) or more drops of primary antibody to cover the tissue completely. Incubate in moist chamber for 30-60 min. b. Rinse with PBS containing 0.05% Tween-20 3 times for 2 minutes each time. 	30-60
5. Reagent 1: HRP Polymer-anti-Goat IgG	<ul style="list-style-type: none"> a. Apply 2 (100 µL) or more drops of HRP Polymer-anti-Goat IgG to cover tissue section and Incubate in moist chamber for 15 min. b. Rinse with PBS containing 0.05% Tween-20 3 times for 2 minutes each time. 	15
6. Reagents 2A, 2B: 2A: DAB Substrate 2B: DAB Chromogen	<ul style="list-style-type: none"> a. Adding 1 drop or 2 drops (for higher contrast) of DAB chromogen concentrate (Reagent 2B) in 1ml of DAB substrate buffer (Reagent 2A). Mix well. b. Apply 2 drops (100 µL) or enough volume of pre-mixed DAB Chromogen to completely cover tissue. Incubate for 5 min. use the prepared DAB solution within 5 hours c. When appropriate color is developed, rinse under tap water gently for about 1-2 minutes. 	5
8. Hematoxylin: Supplied by user.	<ul style="list-style-type: none"> a. Counterstain with 2 (100 ul) or more drops hematoxylin to cover tissue completely and wait about 20 seconds. b. Rinse well with tap water for 1-2 min. c. Put slides in PBS until the color turn blue (about 15-30 seconds.) d. Rinse in distill water, then rinse well with tap water 	20-30 seconds

<p>9. Mounting medium: Supplied by user</p>	<p>Follow the manufacture data sheet procedure for mounting. Recommended product:</p> <ol style="list-style-type: none"> 1. NeoBio Mount AQ: Cat.# NB-00155-3 (18ml), for alcohol soluble substrates (AEC, AP-Red and AP-blue) 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. Can be used with or without cover slip 	<p>Refer to insert</p>
---	--	------------------------

Protocol Notes:

1. The fixation, tissue slide thickness, 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:

DAB may be carcinogenic. Please wear gloves and take other necessary precautions.

Remarks:

For research use only.

Storage:

Store at 4°C.

References:

1. Bisgaard K, Pluzed KP. Use of polymer conjugates in immunohistochemistry: A comparative study of a traditional staining method to a staining method utilizing polymer conjugates. Abstract XXI Intl Cong Intl Acad Pathol and 12th World Cong Acad Environ Pathol. Budapest, Hungary, October 20-25, 1996.

Related products

Product	Catalog No.	Size
NeoStain Poly 1-Step HRP Mouse/Rabbit Bulk kit for DAB	NB-23-00028-1	1L
NeoStain Poly 1-Step HRP Mouse/Rabbit 18ml, 6ml DAB Kit	NB-23-00028-4 / -5	18ml / 6ml
NeoStain Poly 1-Step Mouse Bulk kit for DAB	NB-23-00029-1	110ml
NeoStain Poly 1-Step Mouse 18ml, 6ml DAB Kit	NB-23-00029-4 / -5	18ml / 6ml
NeoStain Poly 1-Step Goat Bulk kit for DAB	NB-23-00031-1	110ml
NeoStain Poly 1-Step Goat 18ml, 6ml DAB Kit	NB-23-00031-2 / -3	18ml / 6ml
NeoStain Poly 1-Step HRP Rat-NM Bulk kit for DAB (no x Mouse)	NB-23-00032-1	110ml
NeoStain Poly 1-Step HRP Rat-NM 18ml, 6ml DAB Kit (no x Mouse)	NB-23-00032-2 / -3	18ml / 6ml
NeoStain Poly 1-Step HRP Mouse-NR Bulk kit for DAB (no x Rat)	NB-23-00033-1	110ml
NeoStain Poly 1-Step HRP Mouse-NR 18ml, 6ml DAB Kit (no x Rat)	NB-23-00033-2 / -3	18ml / 6ml
DAB+ 2 components	NB-23-00148-1	12ml +240ml
NeoBio Mount Perm (Organic)	NB-23-00156	18ml
NeoBio Mount Universal (Aqueous)	NB-23-00157-1 / -2	100ml / 18ml