



Polink-1 HRP Goat AEC Detection System

(Polymer-HRP detection system for AEC staining, biotin-free, Anti-Goat)

Ready-to-use One Step Polymer Detection System

Super Sensitive for AEC Staining

G. 4.00G	Catalog No.	☐ D34-110	110 ml (bulk, w/o chromogen)
Storage: 4-8°C		□ D34-18	18 ml (with AEC, good for 180 slides)
		☐ D34-6	6 ml (with AEC, good for 50 slides)

Intended Use:

Polink-1HRP Goat AEC Detection Kit is designed to use with user supplied goat antibody to detect target antigen on human tissue or cell samples. Polink-1 HRP anti Goat AEC Detection Kit does not cross react with bovine IgG. It is compatible with BSA containing diluent or blocking buffer. Specimen can be frozen or paraffin—embedded tissues, and freshly prepared monolayer cell smears. This detection system is super sensitive when use with AEC chromogen.

Polink-1 1HRP Goat AEC Detection Kit is the ONE step polymer detection system that uses polymeric horseradish peroxidase (HRP) - linked 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.

Kit components:

· ·	5 1 (1)	Reagent 1:	Reagent 2: 2A: substrate buffer (20x)		
Catalog No.	Product Name	Polymer HRP-linked anti goat IgG			
		for AEC	2B: Chromogen (20x)		
		(Ready-to-use)	2C: H ₂ O ₂ (20x)		
D34-110	Polink-1 HRP Goat Bulk for AEC	110ml	Not provided		
D34-60	Polink-1 HRP Goat Bulk for AEC	60ml	Not provided		
D34-18	Polink-1 HRP Goat with AEC kit	18ml	3ml of Reagent 2A		
			6ml of Reagent 2B		
			3ml of Reagent 2C		
D34-6	Polink-1 HRP Goat with AEC kit	6ml	2ml of Reagent 2A		
			4ml of Reagent 2B		
			2ml of Reagent 2C		

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	Staining Procedure	Incubation Time
		(Min.)
Peroxidase Blocking Reagent Supplied by user	a. Incubate slides in peroxidese blocking reagent (Ready-to-use $3\%\ H_2O_2$ solution) for 10 min. b. Rinse the slide using distilled water.	10
HIER Pretreatment: Refer to antibody data sheet.	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. Primary antibody:	Notes: Investigator needs to optimize dilution and incubation times a. Apply 2 (100 μL) or more drops of primary antibody to cover the tissue completely. Incubate in moist chamber for 30-60 min.	30-60

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Supplied by user	b. Rinse with PBS containing 0.05% Tween-20 3 times for 2 minutes each time.	
4. Reagent 1: HRP Polymer-anti-Goat IgG (Ready-to-use)		
5. Reagents 2A, 2B and 2C: AEC Chromogen (20x)	drop of Reagent 2C to 1 mL distilled or deionized water. Mix well. Protect from light and	
6. Hematoxylin:	a. Counterstain with 2 (100 ul) or more drops hematoxylin to cover tissue completely and wait about 20 seconds .	20-30 seconds
Supplied by user.	b. Rinse well with tap water for 1-2 min. c. Put slides in PBS until the color turn blue (about ½ - 1 min.) d. Rinse in distill water, then rinse well with tap water	
7. Mounting medium:	Mounting medium: Follow the manufacture data sheet procedure for mounting. Recommended product:	
Supplied by user	1. GB-Mount: Cat. No. E01-18 (18ml), for alcohol soluble substrates (AEC, AP-Red and AP-blue) 2. O-Mount: Cat. No. E02-18 (18ml), for DAB and BCIP/NBT 3. Simpo-Mount: Cat.No. E03-18 (18ml), or E03-100 (100ml), universal permanent mounting medium. Can be used with or without cover slip	

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.

Related Products:

Product	Catalog No.	Size	Size Product		Catalog No.	Size
Polink-1 HRP Mouse Bulk kit for AEC	D15-110	110ml		*Polink-1 HRP Rat-NM 18ml, 6ml	D36-18 / D36-6	18ml / 6ml
				Kit		
Polink-1 HRP Mouse 18ml, 6ml AEC Kit	D15-18 / D15-6	18ml / 6ml		**Polink-1 HRP Mouse-NR Bulk kit	D56-110	110ml
				for AEC		
Polink-1 HRP Rabbit Bulk kit for AEC	D16-110	110ml		**Polink-1 HRP Mouse-NR 18ml,	D56-18 / D56-6	18ml / 6ml
				6ml AEC Kit		
Polink-1 HRP Rabbit 18ml, 6ml AEC Kit	D16-18 / D16-6	18ml / 6ml		AEC Kit	C01-12	12ml
Polink-1 HRP Broad Bulk kit for AEC	D14-110	110ml		GB-Mount (Aqueous)	E01-18	18ml
Polink-1 HRP Broad 18ml, 6ml AEC Kit	D14-18 / D14-6	18ml / 6ml		Simpo-Mount (Aqueous)	E03-100 /E03-18	100ml / 18ml
*Polink-1 HRP Rat-NM Bulk kit for AEC	D36-110	110ml				

^{*}Polink-1 HRP Rat-NM kit does not cross react with mouse primary antibody
**Polink -1 HRP Mouse-NR kit does not cross react with Rat primary antibody

Precautious:

Please wear gloves and take other necessary precautions.

Remarks:

For research use only.

References:

- 1. <u>Bisgaard K, Pluzed KP</u>. Use of polymer conjugates in immunohitochemistry: A comparative study of a traditional staining method to a staining method utilizing polymer conjugates. <u>Abstract</u> XXI Intl Cong Intl Acad Pathol and 12th World Cong Acad Environ Pathol. Budapest, Hungry, October 20-25, 1996.
- 2. Shi ZR. Itzkowitz SH, Kim YS. A comparison of three immunoperoxidase techniques for antigen detection in colorectal carcimoma tissues. J Hitochem Cytochem 36:317-322,