

BrightVision Poly-AP-Anti Ms/Rb/Rt IgG
Biotin-free, One Component
Ready-to-use

Application:

BrightVision Histostaining reagents utilize a novel controlled polymerisation technology to prepare polymeric AP-linker antibody conjugates. Comparing to conventional biotin-streptavidin based detection kits, BrightVision histostaining kits have the advantages of higher amplification power, biotin free and more consistent immunostaining outcomes on archival tissues and on difficult-to-work antibodies. (ref-1). These advantages would bring to laboratories the benefit of more accurate result, faster turn-around, less trouble shooting and better costs-saving.

Kit Components: Poly-AP-Anti-mouse/rabbit/rat IgG (ready-to-use).

Availability:

Catalog No.	Contents	Volume
DPVO999AP	Poly-AP-Anti-mouse/rabbit/rat IgG (ready-to-use)	1000 ml
DPVO500AP	Poly-AP-Anti-mouse/rabbit/rat IgG (ready-to-use)	500 ml
DPVO110AP	Poly-AP-Anti-mouse/rabbit/rat IgG (ready-to-use)	110 ml
DPVO55AP	Poly-AP-Anti-mouse/rabbit/rat IgG (ready-to-use)	55 ml

Species of origin: Goat

Antigen Specificity: Anti-Mouse IgG (H+L), Anti-Rabbit IgG (H+L), Anti-Rat IgG (H+L)

Enzyme Conjugate: Alkaline Phosphatase






Recommended Staining Protocol (kit components in bold):

1. Deparaffinize and rehydrate tissue section.
2. To reduce non-specific background staining due to endogenous peroxidase, incubate slide in hydrogen peroxide for 10-15 minutes.
3. Wash 2 times in PBS or TBS wash buffer.
4. If required, incubate tissue in digestive enzyme (or appropriate pre-treatment).
5. Wash 2 times in PBS or TBS wash buffer.
6. (Optional) Apply Pre-antibody Blocking Solution (NGS) and incubate for 5 minutes at room temperature to block non-specific background staining.
Note: Do not exceed 10 minutes or there may be a reduction in desired stain.
7. Wash 2 times in PBS or TBS wash buffer.
8. Apply primary mouse or rabbit antibody and incubate according to manufacturer's protocol.
9. Wash 2 times in PBS or TBS wash buffer.
10. Apply **Poly-AP-Goat anti Mouse/Rabbit/Rat IgG**, and incubate for 30 minutes at room temperature.
11. Wash 2 times in PBS or TBS wash buffer.
12. Incubate with Incubate with Fast-Red or New-Fuchsin solution.
13. Counterstain and coverslip.

Note: Pre-antibody blocking is optional and can be omitted if primary antibodies are diluted in buffers containing 2-10% normal goat serum.

Reference:

- 1) Shan-Rong Shi, James Guo, Richard J. cote, Lillian Young, Debra Hawes, Yan Shi, Sandra Thu and Clive R. Taylor, Applied Immunohistochemistry & Molecular Morphology, vol 7, 201-208, 1999.

	CE Mark (European Union Countries)		In Vitro Diagnostic Medical Device		Immunologic bv Typograaf 16 6921 VB Duiven The Netherlands
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