

## UltraVision LP Detection System AP Polymer & Fast Red Chromogen

### DESCRIPTION:

**UltraVision LP** is the latest technology in polymeric labeling. Polymer detection methods have been shown to provide increased sensitivity and detection simplicity. This second-generation polymer system is composed of smaller polymer subunits that minimize conflicts in binding the target protein. Decreased binding conflicts result in more consistent staining and better signal amplification.<sup>1</sup> Ultimately, this gives the user higher sensitivity and antibody efficiency.<sup>2</sup> With **UltraVision LP**, you use less antibody and obtain better signal-to-noise ratios. **UltraVision LP** is also biotin-free, which eliminates background staining found with traditional biotin-based detection methods.

**SPECIFICITY:** Anti-Mouse IgG (H+L), Anti-Rabbit IgG (H+L)  
**ENZYME:** Alkaline Phosphatase  
**CHROMOGEN/SUBSTRATE:** Fast Red/Naphthol Phosphate Buffer

**AVAILABILITY:**

<u>Catalog #</u>	<u>Slide Volume</u>
TL-012-AF	120-240 slides
TL-015-AF	150-300 slides

### STAINING PROTOCOL (kit components in bold):

1. Deparaffinize and rehydrate tissue section.
2. Wash 2 times in buffer.
3. If required, incubate tissue in digestive enzyme (or appropriate pretreatment).
4. Wash 4 times in buffer.
5. To reduce nonspecific background staining due to endogenous peroxidase, incubate slide in **Hydrogen Peroxide Block** for 10-15 minutes.
6. Wash 4 times in buffer.
7. Apply **Ultra V Block** and incubate for 5 minutes at room temperature to block nonspecific background staining. Note: Do not exceed 10 minutes or there may be a reduction in desired stain. (May be omitted if primary antibodies are diluted in buffers containing 5-10% normal goat serum.)
8. Wash (Optional).
9. Apply primary antibody and incubate according to manufacturer's recommended protocol.
10. Wash 4 times in buffer.

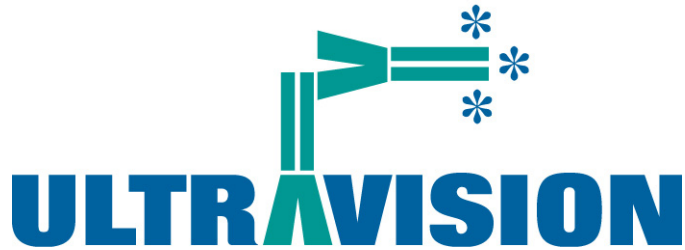
<sup>1</sup> Shan-Rong Shi, James Guo, Richard J. Cote, Lillian Young, Debra Hawes, Yan Shi, Sandra Thu, and Clive R. Taylor, Applied Immunohistochemistry & Molecular Morphology, vol 7, 201-208, 1999.

<sup>2</sup> Karen Petrosyan, Rosalba Tamayo, and Daisy Joseph, "Sensitivity of a Novel Biotin-free Detection Reagent (PowerVision+) for Immunohistochemistry" J. Histotechnology, vol 25, 247-250, 2002.



## SPECIFICATION SHEET

11. Apply **Primary Antibody Enhancer** and incubate for 20 min at room temperature.
12. Wash 4 times in buffer.
13. Incubate with **AP Polymer** anti-Mouse/Rabbit IgG for 30 min at room temperature.
14. Wash 4 times in buffer.
15. Add 1 **Fast Red tablet** to 5 ml of **Naphthol Phosphate Substrate** and shake until tablet is dissolved. Apply to tissue section and incubate for 10-20 minutes or until desired stain intensity is achieved.
16. Wash 4 times in DI water.
17. Counterstain and coverslip using a permanent mounting media.



**LIMITATIONS:**

This product is available for laboratory use only - not for therapeutic work. LabVision Corporation will not be held responsible for patent infringement or other violation that may occur with the use of this product.

**STORAGE:** Store at 4-6°C. Each component is stable for 18 months.

**TROUBLESHOOTING GUIDE**

**OVERSTAINING:**

1. Concentration of the primary antibody was too high or the incubation time was too long.
2. Temperature during incubation was too high.
3. Incubation time with label was too long.

**NONSPECIFIC BACKGROUND STAINING:**

1. Rinsing between steps was inadequate.
2. Tissue was allowed to dry with reagents on.
3. Folds in tissue trapped reagents.
4. Tissue contains endogenous peroxidase.
5. Antigen migrated in tissue.
6. Excessive tissue adhesive on slides.
7. Inadequate blocking with protein block.

**WEAK STAINING:**

1. Primary antibody concentration was too low or incubation time was too short.
2. Reagents have degraded.
3. Inadequate removal of wash buffer between steps, resulting in dilution of reagents.
4. Counterstain or mounting media were incompatible and dissolved the chromogen reaction product.
5. Room temperature was excessively cool.
6. The primary antibody does not recognize an antigen that survives fixation and embedding in high enough amounts.
7. Excessive incubation with protein block (Ultra Block or normal serum).

**NO STAINING:**

1. Steps were inadvertently left out.



SPECIFICATION SHEET

2. There is no antigen in the tissue.
3. The primary antibody is not compatible with the detection method.
4. Chromogenic substrate has been replaced with another that is not intended for use with peroxidase.
5. One or more components of the kit have been inactivated.