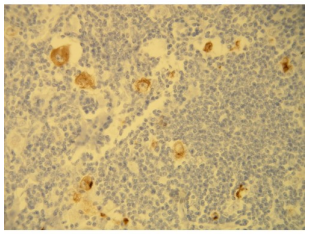
Epstein-Barr Virus Latent Membrane Protein 1 (EBV LMP1) Antibody

Catalogue No.:abx227096



IHC-P analysis of HRS classical Hodgkin Lymphoma cells using EBV LMP1 antibody, showing cytoplasmic expression of the EBV LMP-1 protein (4 µm section).

Epstein-Barr Virus Latent Membrane Protein 1 (EBV LMP1) Antibody is a Rabbit Monoclonal antibody for the detection of EBV/LMP-1.

Target:	Epstein-Barr Virus Latent Membrane Protein 1 (EBV LMP1)
Clonality:	Monoclonal
Clone:	W978
Reactivity:	Human
Tested Applications:	IHC
Host:	Rabbit
Recommended dilutions:	IHC-P: 1/100. Optimal dilutions/concentrations should be determined by the end user.
Conjugation:	Unconjugated
Immunogen:	Synthetic peptide derived from the C-terminal region of Epstein-Barr virus, Latent Membrane Protein-1 (Ser369-Tyr384).
Isotype:	IgG
Form:	Liquid
Purification:	Purified from rabbit antiserum by proprietary techniques.
Storage:	Store at 2-8°C.
UniProt Primary AC:	P03230 (UniProt, ExPASy)

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Buffer: 20 mM Tris-HCl, pH 8.0, containing 20 mg/ml BSA and 0.05% NaN3.

Concentration: 0.5 mg/ml

Note: THIS PRODUCT IS FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC, THERAPEUTIC OR COSMETIC PROCEDURES. NOT FOR HUMAN OR ANIMAL CONSUMPTION.

Directions for use:

Suggested IHC-P Protocol

1. Preparation of Citrate Buffer (10 mM Citric Acid, 0.05% Tween-20, pH 6.0): Dissolve 1.92 g of anhydrous citric acid in 700 ml of distilled water. Adjust pH to 6.0 with 1 M NaOH and then add 0.5 ml of Tween-20 and mix thoroughly. Adjust the final volume to 1 L with distilled water. Store this solution at room temperature for up to 3 months or at 4 °C for long-term storage.
2. Preparation of Wash Buffer: Use 0.05 M Tris-HCl, pH 7.6, containing 0.2% Tween-20.
3. Deparaffinize the section in 3 changes of xylene, 5 minutes each.
4. Wash the section in 96%, 80% and 70% ethanol, 10 minutes each.
5. Rinse in distilled water.
6. Block endogenous peroxidase by incubating the tissue in 3% hydrogen peroxide (H_2O_2) for 10 minutes.
7. Wash in distilled water for 5 minutes.
8. Antigen retrieval: immerse the slide in Citrate buffer, pH 6.0, and incubate in a water bath for 40 minutes at 96 °C.
9. Remove the slide from the water bath and allow to stand at room temperature (in Citrate buffer, pH 6.0) for 20 minutes.
10. Rinse in distilled water.
11. Wash in Wash Buffer for 5 minutes.
12. Incubate the section with primary antibody at 1/100 - 1/200 dilution for 1 hour in a closed wet chamber. It is recommended to use abx291502 Primary Antibody Diluent or a diluent containing protease-free BSA (> 1 mg/ml) to dilute this antibody.
13. Wash twice with Wash Buffer, 5 minutes each.
14. Add the secondary antibody and proceed to standard immunohistochemistry protocol (HRP - Peroxide - DAB). It is recommended to use abx291501 Rabbit and Mouse HRP/DAB Detection Kit.
15. Wash twice in Wash Buffer, 5 minutes each.
16. Apply the DAB chromagen for 1-3 minutes.
17. Rinse in water for 10 minutes.
18. Stain in hematoxylin for 5 minutes.
19. Wash in water for 10 minutes.
20. Dehydrate the section in 2 changes of 96% ethanol, 5 minutes each.
21. Wash the section in 2 changes of xylene, 2 minutes each.
22. Mount the slide for observation.

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