

Product Datasheet

Bcl-2 Antibody (P21-A)

NBP1-79011-100ul

Unit Size: 100 ul

Store at 4C. Do not freeze.

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NBP1-79011-100ul

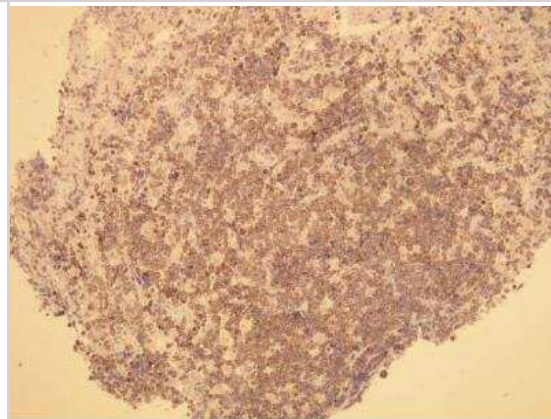
Bcl-2 Antibody (P21-A)

Product Information	
Unit Size	100 ul
Concentration	Please see the vial label for concentration. If unlisted please contact technical services.
Storage	Store at 4C. Do not freeze.
Clonality	Monoclonal
Clone	P21-A
Preservative	0.05% Sodium Azide
Isotype	IgG
Purity	Immunogen affinity purified
Buffer	20mM Tris/HCl (pH 8.0) and 20 mg/ml BSA
Product Description	
Host	Rabbit
Gene ID	596
Gene Symbol	BCL2
Species	Human
Immunogen	peptide derived from the sequence close to N-terminal region of human bcl-2
Notes	This antibody is immunoaffinity purified with immunogenic peptide as a ligand.
Product Application Details	
Applications	Immunohistochemistry, Immunohistochemistry-Paraffin
Recommended Dilutions	Immunohistochemistry 1:10-1:500, Immunohistochemistry-Paraffin 1:100-1:200

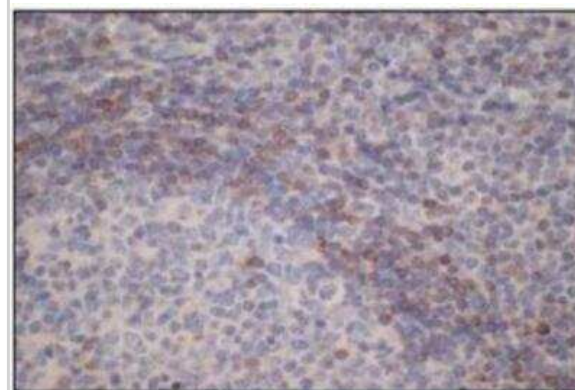


Images

Immunohistochemistry-Paraffin: Bcl-2 Antibody (P21-A) [NBP1-79011] - Expression in extranodal marginal zone lymphoma of mucosa - associated lymphoid tissue (MALT lymphoma) of the stomach.



Immunohistochemistry-Paraffin: Bcl-2 Antibody (P21-A) [NBP1-79011] - Formalin-fixed and paraffin-embedded human tonsil tissue (4 um) stained with anti - Bcl-2 antibody shows strong membrane immunostaining of follicular cells.



Immunohistochemistry-Paraffin: Bcl-2 Antibody (P21-A) [NBP1-79011] - Expression in diffuse large B-cell lymphoma.



Procedures

Immunohistochemistry-Paraffin protocol for Bcl-2 Antibody (NBP1-79011)

IHC-P Protocol (NBP1-79011):

1. Deparaffinize the section in 3 changes of xylene, 5 minutes each.
2. Wash the section in 96%, 80% and 70% benzyl alcohol for 5 minutes each.
3. Rinse in distilled water.
4. Block the endogenous peroxidase by incubating the tissue in 3% hydrogen peroxide (H₂O₂) for 10 minutes.
5. Wash in distilled water.
6. For antigen retrieval: immerse the slide in Tris-EDTA buffer, pH 9.0, 0.05% Tween- 20*, and incubate at 90C in water bath for 30 minutes. (Alternatively adjust to your own protocol, keeping the required pH)
7. Remove the staining to room temperature and let the slide to cool (in TRIS-EDTA buffer, pH 9.0) for 15 minutes.
8. Rinse in distilled water.
9. Wash in 0.05 M Tris-HCl , pH 7.6 buffer supplemented with 0.5% of Tween-20 (buffer A) for 5 minutes.
10. Incubate the section with primary antibody diluted in buffer A at the dilution 1:100 - 200 for 1 hour in the closed wet chamber.
11. Wash twice 5 minutes with buffer A.
12. Apply the secondary antibody (the protocol depends on the supplier), and proceed to standard immunohistochemistry protocol (HRP - Peroxide - DAB).
13. Wash twice 5 minutes with buffer A.
14. Apply the chromogen (DAB), 10 minutes.
15. Wash in water - 10 minutes.
16. Stain in hematoxylin for 5 minutes.
17. Wash in water - 10 minutes.
18. Dehydrate the section in 2 changes of 96% benzyl alcohol for 5 minutes each.
19. Wash the section in 2 changes of xylene for 2 minutes each.
20. Mount the slide for observation



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Limitations

This product is for research use only and is not approved for use in humans or in clinical diagnosis. Primary Antibodies are guaranteed for 1 year from date of receipt.

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