

bcl-6

Availability/Contents:	Item #	Volume
	A20040	2 ml
	A00040	6 ml
	A00040.25	25 ml

Description:


Species:	Rabbit
Immunogen:	Synthetic peptide corresponding to amino acids 687-706 mapping at the carboxy terminus of bcl-6 of human origin.
Clone:	Polyclonal
Isotype:	N/A
Format:	This antibody has been pretitered and quality controlled to work on formalin-fixed paraffin-embedded as well as acetone fixed cryostat tissue sections. No further titration is required.
Specificity:	Relatively little is known regarding the molecular pathogenesis of diffuse large-cell lymphoma (DLCL). DLCL accounts for approximately 40% of all non Hodgkin lymphomas. Chromosomal translocations involving reciprocal recombination between band 3q27 and several other chromosomal sites are involved in 8 to 12% of non-Hodgkin's lymphomas; a breakpoint cluster region at 3q27 has been identified and designated bcl-6. More recently this gene has been shown to encode a protein of 706 amino acids with predicted molecular weight of 79 kD. This bcl-6 protein has homology with zinc finger-transcription factors and was found to be rearranged in 13 of 39 DLCL samples.
Species Reactivity:	Human, Mouse, Rat.
Positive Control:	Tonsil
Cellular Localization:	Nuclear
Titer/Working Dilution:	No further dilution is required.
Microbiological State:	This product is not sterile.


Uses/Limitations: For In Vitro Diagnostic Use.
This product is intended for qualitative immunohistochemistry with normal and neoplastic formalin-fixed, paraffin-embedded tissue sections, to be viewed by light microscopy.
Do not use past expiration date.




Storage and Stability: 2-8° Centigrade.
Product is stable for 24 months from date of manufacture.
If reagent is not stored as recommended, performance must be validated by the user.

Procedure:

1. Tissue Section Pretreatment: Staining of formalin fixed, paraffin embedded tissue sections is enhanced by pretreatment with Citrate Plus (ScyTek catalog# CPL500) or 10mM citrate buffer, pH 6.0 (ScyTek Catalog# CBB500, see IFU for instructions).
2. Primary Antibody Incubation Time: We suggest an incubation period of 30-60 minutes at room temperature. However, depending upon the fixation conditions and the staining system employed, optimal incubation should

Storage: 2°C  8°C

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be determined by the user.

3. Visualization: For maximum staining intensity we recommend the "Retrieval HRP Anti-Polyvalent Lab Pack" (ScyTek catalog# RPL125, see IFU for instructions) combined with the "DAB Chromogen/Substrate Bulk Pack (High Contrast)" (ScyTek catalog# ACV500, see IFU for instructions).


Precautions: Contains Sodium Azide as a preservative (0.09% w/v).
Do not pipette by mouth.
Avoid contact of reagents and specimens with skin and mucous membranes.
Avoid microbial contamination of reagents or increased nonspecific staining may occur.
This product contains no hazardous material at a reportable concentration according to U.S. 29 CFR 1910.1200, OSHA Hazardous Communication Standard and EC Directive 91/155/EC.


References:




1. Baron et al. Proc Natl Acad Sci (USA) 90: 5262, 1993
2. Bastard et al. Blood 79: 2527, 1992.
3. Ye et al. Science 262: 747, 1993.
4. Ye et al. Cancer Res 53: 2732, 1993

Warranty:

No products or "Instructions For Use (IFU)" are to be construed as a recommendation for use in violation of any patents. We make no representations, warranties or assurances as to the accuracy or completeness of information provided on our IFU or website. Our warranty is limited to the actual price paid for the product. ScyTek Laboratories, Inc. is not liable for any property damage, personal injury, time or effort or economic loss caused by our products. Immunohistochemistry is a complex technique involving both histological and immunological detection methods. Tissue processing and handling prior to immunostaining can cause inconsistent results. Variations in fixation and embedding or the inherent nature of the tissue specimen may cause variations in results. Endogenous peroxidase activity or pseudoperoxidase activity in erythrocytes and endogenous biotin may cause non-specific staining depending on detection system used.

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