abcam

Product datasheet

Anti-Lambda Light chain antibody [EPR5367-62] - BSA and Azide free ab185131

אילשעבע RabMAb

画像数6

製品の概要			
製品名	Anti-Lambda Light chain antibody [EPR5367-62] - BSA and Azide free		
製品の詳細	Rabbit monoclonal [EPR5367-62] to Lambda Light chain - BSA and Azide free		
由来種	Rabbit		
アプリケーション	適用あり: IHC-P, ICC/IF, Flow Cyt (Intra), ELISA, WB		
種交差性	交差種: Human		
免疫原	Full length native protein (purified) corresponding to Human Lambda Light chain.		
特記事項	ab185131 is the carrier-free version of ab124719 .		
	Our <u>carrier-free</u> antibodies are typically supplied in a PBS-only formulation, purified and free of BSA, sodium azide and glycerol. The carrier-free buffer and high concentration allow for increased conjugation efficiency.		
	This conjugation-ready format is designed for use with fluorochromes, metal isotopes, oligonucleotides, and enzymes, which makes them ideal for antibody labelling, functional and cell-based assays, flow-based assays (e.g. mass cytometry) and Multiplex Imaging applications.		
	Use our conjugation kits for antibody conjugates that are ready-to-use in as little as 20 minutes with <1 minute hands-on-time and 100% antibody recovery: available for fluorescent dyes, HRP, biotin and gold.		
	This product is compatible with the Maxpar [®] Antibody Labeling Kit from Fluidigm, without the need for antibody preparation. Maxpar [®] is a trademark of Fluidigm Canada Inc.		
	 This product is a recombinant monoclonal antibody, which offers several advantages including: High batch-to-batch consistency and reproducibility Improved sensitivity and specificity Long-term security of supply Animal-free production For more information <u>see here</u>. Our RabMAb[®] technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to <u>RabMAb[®] patents</u>. 		
	Mouse, Rat: We have preliminary internal testing data to indicate this antibody may not react with these species. Please contact us for more information.		

製品の特性			
製品の状態	Liquid		
保存方法	Shipped at 4°C. Store at +4°C. Do Not Freeze.		
バッファー	pH: 7.2 Constituent: PBS		
キャリア・フリー	はい		
精製度	Protein A purified		
ポリ/モノ	モノクローナル		
クローン名	EPR5367-62		
アイソタイプ	lgG		

アプリケーション

The Abpromise guarantee

<u>Abpromise保証は、</u>次のテスト済みアプリケーションにおけるab185131の使用に適用されます

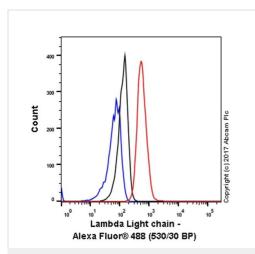
アプリケーションノートには、推奨の開始希釈率がありますが、適切な希釈率につきましてはご検討ください。

アプリケーション	Abreviews	特記事項
IHC-P		Use at an assay dependent concentration. Perform heat mediated antigen retrieval before commencing with IHC staining protocol. See IHC antigen retrieval protocols .
ICC/IF		Use at an assay dependent concentration.
Flow Cyt (Intra)		Use at an assay dependent concentration.
ELISA		Use at an assay dependent concentration.
WB		Use at an assay dependent concentration. Predicted molecular weight: 25 kDa.

ターゲット情報

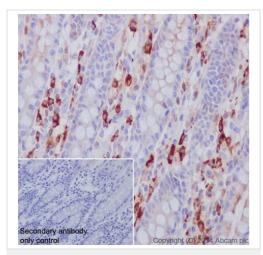
関連性

All five immunoglobulin classes share the same basic four polypeptide chain structure of two heavy-chains and two light chains. There are five heavy chain types, and two light-chain types (Kappa and Lambda) both having a molecular weight of 22.5kDa. Any heavy-chain type can associate with either light-chain type, but on any immunoglobulin molecule both light-chains are of the same type. Kappa and Lambda consist of a variable region and a constant region and can easily be differentiated by the antigenic properties of the constant region. The ratio of Kappa to Lambda is 70:30, the vast majority of which is bound to heavy-chain in immunoglobulin. In normal individuals low levels of free light-chain arepresent in serum (kappa, 1.6-15.2 mg/L; Lambda, 0.4-4.2mg/L), with the occurrence of multiple myeloma or other B-cell malignancies these levels can be greatly elevated and can be found at high levels in the urine (Bence-Jones proteins).



Flow Cytometry (Intracellular) - Anti-Lambda Light chain antibody [EPR5367-62] - BSA and Azide free (ab185131) Intracellular Flow Cytometry analysis of Ramos (human Burkitt's lymphoma) cells labeling Lambda Light chain with unpurified **ab124719** at 1/50 dilution (10ug/ml) (red). Cells were fixed with 4% paraformaldehyde and permeabilised with 90% methanol. A Goat anti rabbit IgG (Alexa Fluor[®] 488) (1/2000 dilution) was used as the secondary antibody. Rabbit monoclonal IgG (Black) was used as the isotype control, cells without incubation with primary antibody and secondary antibody (Blue) were used as the unlabeled control.

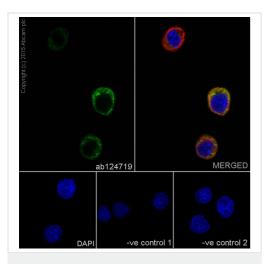
This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab124719</u>).



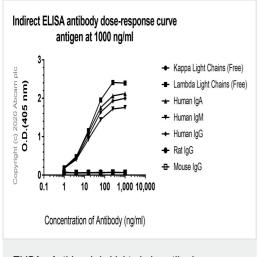
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Lambda Light chain antibody [EPR5367-62] - BSA and Azide free (ab185131)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human colon tissue labelling Lambda Light chain with purified **ab124719** at 1/300. Heat mediated antigen retrieval was performed using Tris/EDTA buffer pH 9. **ab97051**, a HRP-conjugated goat anti-rabbit IgG (H+L) was used as the secondary antibody (1/500). Negative control using PBS instead of primary antibody. Counterstained with hematoxylin.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (**ab124719**).



Immunocytochemistry/ Immunofluorescence - Anti-Lambda Light chain antibody [EPR5367-62] - BSA and Azide free (ab185131)



ELISA - Anti-Lambda Light chain antibody [EPR5367-62] - BSA and Azide free (ab185131) Immunocytochemistry/Immunofluorescence analysis of Ramos cells labelling Lambda Light chain with purified <u>ab124719</u> at 1/250. Cells were fixed with 4% paraformaldehyde and permeabilized with 0.1% Triton X-100. <u>ab150077</u>, an Alexa Fluor[®] 488-conjugated goat antirabbit IgG (1/500) was used as the secondary antibody. DAPI (blue) was used as the nuclear counterstain. <u>ab7291</u>, a mouse anti-tubulin (1/1000) and <u>ab150120</u>, an Alexa Fluor[®] 594-conjugated goat antimouse IgG (1/500) were also used.

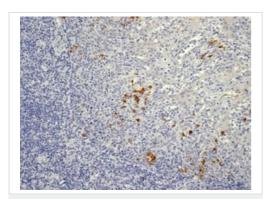
Control 1: primary antibody (1/250) and secondary antibody, <u>ab150120</u>, an Alexa Fluor[®] 594-conjugated goat anti-mouse IgG (1/500).

Control 2: <u>**ab7291**</u> (1/1000) and secondary antibody, <u>**ab150077**</u>, an Alexa Fluor[®] 488-conjugated goat anti-rabbit lgG (1/500).

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab124719</u>).

ELISA analysis of Human Kappa light chain (Free), Human Lambda Light Chains (Free), Human IgA, Human IgM, Human IgG, Rat IgG, Mouse IgG at 1000 ng/mL with <u>ab124719</u> at 1000~0ng/mL. An Alkaline Phosphatase-conjugated AffiniPure Goat Anti-Rabbit IgG (H+L) at 1/2500 dilution was used as the secondary antibody.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab124719</u>).



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Lambda Light chain antibody [EPR5367-62] - BSA and Azide free (ab185131)



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Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human tonsil tissue labelling Lambda Light chain with unpurified **ab124719** at a dilution of 1/250.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab124719</u>).

Perform heat mediated antigen retrieval before commencing with IHC staining protocol.

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