AB-PK540 Bcr-pY644 Antibody

Phosphosite-specific polyclonal antibody for monitoring the phosphorylation of human protein-serine/threonine kinase Bcr



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Name Long:	Breakpoint cluster region protein
Alias:	ALL; BCR; BCR1; Breakpoint cluster region; CML; D22S11; D22S662; PHL; Renal carcinoma antigen NY-REN-26; FLJ16453; CCDS13806.1; Q12843; Q12845; Q12848; Q14020; Q13846; ENSG00000186716
UniProt ID:	P11274
Sequence Predicted Mass (KDa):	142.819 (1271 AA; P11274); 137.729 (1227 AA; P11274-2)
Observed SDS-PAGE Mass (KDa):	150-170
Immunogen	
Antibody Immunogen Source:	Human Bcr sequence peptide Cat. No.: PE-04AIO99
Antibody Immunogen Sequence:	TLL(pY)KPV(bA)C (bA) = beta-alanine
Location in Target:	Corresponds to amino acid residues T641 to V647; In the Rho-GEF domain. One of the main in vivo phosphorylation sites in Bcr.
Peptide Type:	For phosphosite-specific recognition of target.
Target Phosphosite:	Tyr-644
Production Antibody Host Species:	Rabbit
Antibody Type:	Polyclonal
Antibody Ig Isotype Clone Lot:	Immunoglobulin G
	The immunizing peptide was produced by solid phase synthesis on a multipep
Production Method:	sera from each animal was applied onto an agarose column to which the immunogen peptide was thio-linked. Antibody was eluted from the column with
	was assessed by analytical hplc and the amino acid sequence confirmed by mass spectrometry analysis. This peptide was coupled to KLH prior to immunization into rabbits. New Zealand White rabbits were subcutaneously injected with KLH-coupled immunizing peptide every 4 weeks for 4 months. The sera from each animal was applied onto an agarose column to which the immunogen peptide was thio-linked. Antibody was eluted from the column with 0.1 M glycine, pH 2.5. Subsequently, the antibody solution was neutralized to pH 7.0 with saturated Tris. This antibody was also subject to negative purification
Antibody Amount:	was assessed by analytical hplc and the amino acid sequence confirmed by mass spectrometry analysis. This peptide was coupled to KLH prior to immunization into rabbits. New Zealand White rabbits were subcutaneously injected with KLH-coupled immunizing peptide every 4 weeks for 4 months. The sera from each animal was applied onto an agarose column to which the immunogen peptide was thio-linked. Antibody was eluted from the column with 0.1 M glycine, pH 2.5. Subsequently, the antibody solution was neutralized to pH 7.0 with saturated Tris. This antibody was also subject to negative purification over phosphotyrosine-agarose.
Antibody Amount: Antibody Concentration:	was assessed by analytical hplc and the amino acid sequence confirmed by mass spectrometry analysis. This peptide was coupled to KLH prior to immunization into rabbits. New Zealand White rabbits were subcutaneously injected with KLH-coupled immunizing peptide every 4 weeks for 4 months. The sera from each animal was applied onto an agarose column to which the immunogen peptide was thio-linked. Antibody was eluted from the column with 0.1 M glycine, pH 2.5. Subsequently, the antibody solution was neutralized to pH 7.0 with saturated Tris. This antibody was also subject to negative purification over phosphotyrosine-agarose.
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Applications	
Product Use:	Western blotting Antibody microarrays
Antibody Dilution Recommended:	2 µg/ml for immunoblotting
Antibody Species Reactivity:	Human, mouse, rat and many other mammals
Antibody Positive Controls:	Very strong immunoreactivity with immunogen peptide on dot blots.
Overall Antibody Specificity:	Very high selectivity
Antibody Cross Reactivities:	No cross-reactivities with other proteins in A431 and Jurkat cells, and sea star oocytes, but a weak 48 KDa cross-reactive protein was detected in A431 cells.

This product is for in vitro research use only and is not intended for use in humans or animals.

For more information on our products please visit <u>www.kinexusproducts.ca</u> or contact us at 1-866-KINEXUS(546-3987)