AB-NK205-3 ARaf-2 Antibody



Pan-specific polyclonal antibody for monitoring the expression of human protein-serine/threonine kinase Raf-A (ARaf)

Canada V6P 6T3

Address: 8755 Ash Street, Suite 1 Vancouver, British Columbia, Email: info@kinexus.ca Phone: 604-323-2547

Target Protein	
Name Long:	A-Raf proto-oncogene serine/threonine-protein kinase
Alias:	ARAF; A-RAF; ARAF1; A-raf-1; Kinase A-Raf; PKS; PKS2; Proto-oncogene Pks; V-raf murine sarcoma 3611 viral oncogene; RP1-230G1.1-005; RP1-230G1_1; A6NIT1; B4DMG5; Q5H9B2; Q96II5ENSG00000078061
UniProt ID:	P10398
Sequence Predicted Mass (KDa):	67.585 (606 AA; P10398); 20.904 (186 AA; P10398-2)
Observed SDS-PAGE Mass (KDa):	63-70
Immunogen	
Antibody Immunogen Source:	Human Raf-A (ARaf) sequence peptide Cat. No.: PE-01ASF80
Antibody Immunogen Sequence:	CGRKTVTAWDTAIAPL
Location in Target:	Corresponds to amino acid residues G67 to L81;
Peptide Type:	For pan-specific recognition of target expression levels.
Target Phosphosite:	Not phosphorylated
Production	
Antibody Host Species:	Rabbit
Antibody Type:	Polyclonal
	Polyclonal Immunoglobulin G
Antibody Type:	Polyclonal
Antibody Type: Antibody Ig Isotype Clone Lot:	Polyclonal Immunoglobulin G The immunizing peptide was produced by solid phase synthesis on a multipep peptide synthesizer and purified by reverse-phase hplc chromatography. Purity 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
Antibody Type: Antibody Ig Isotype Clone Lot: Production Method:	Polyclonal Immunoglobulin G The immunizing peptide was produced by solid phase synthesis on a multipep peptide synthesizer and purified by reverse-phase hplc chromatography. Purity 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.
Antibody Type: Antibody Ig Isotype Clone Lot: Production Method: Antibody Amount:	Polyclonal Immunoglobulin G The immunizing peptide was produced by solid phase synthesis on a multipep peptide synthesizer and purified by reverse-phase hplc chromatography. Purity 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. 25 µg
Antibody Type: Antibody Ig Isotype Clone Lot: Production Method: Antibody Amount: Antibody Concentration:	PolyclonalImmunoglobulin GThe immunizing peptide was produced by solid phase synthesis on a multipep peptide synthesizer and purified by reverse-phase hplc chromatography. Purity 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.25 μg1 mg/ml





Address: 8755 Ash Street, Suite 1 Vancouver, British Columbia, Canada V6P 6T3

Email: info@kinexus.ca Phone: 604-323-2547

Applications	
Product Use:	Western blotting Antibody microarrays
Antibody Dilution Recommended:	1 µg/ml for immunoblotting
Antibody Species Reactivity:	Human, mouse, rat and many other mammals
Overall Antibody Specificity:	High selectivity
Antibody Cross Reactivities:	No immunoreactivity on protein dot blots with recombinant human B-Raf and Raf1.

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