For Research Use Only

## TFG Polyclonal antibody Catalog Number: 11571-1-AP Featured Produ

Featured Product



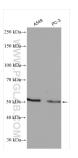


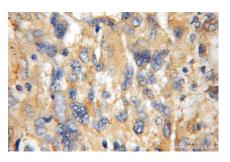
Basic Information	Catalog Number: 11571-1-AP	GenBank Accession Number: BC023599	Purification Method: Antigen affinity purification
	Size:	GenelD (NCBI):	Recommended Dilutions:
	500 µg/ml	10342	WB 1:500-1:2000
	Source: Rabbit	UNIPROT ID: Q92734	IHC 1:20-1:200 IF/ICC 1:50-1:500
	Isotype: IgG	Full Name: TRK-fused gene	
	Immunogen Catalog Number: AG2151	Calculated MW: 400 aa, 43 kDa	
		Observed MW: 50-55 kDa	
Applications	Tested Applications:	Positive Controls:	
	WB, IHC, IF/ICC, ELISA Cited Applications:	WB : A549 cells, PC-3 cells	
	WB, IF		ıman gliomas tissue,
	Species Specificity: human, mouse, rat	IF/ICC	A 549 cells,
	Cited Species: human, mouse		
	Note-IHC: suggested antigen retrieval with TE buffer pH 9.0; (*) Alternatively, antigen retrieval may be performed with citrate buffer pH 6.0		
	Protein TFG (TRK-fused gene protein) plays a role in regulating phosphotyrosine-specific phosphatase-1 activity. Mutations in TFG may have important clinical relevance for current therapeutic strategies to treat metastatic melanoma. Defects in TFG are a cause of thyroid papillary carcinoma (TPC), a common tumor of the thyroid that typically arises as an irregular, solid or cystic mass from otherwise normal thyroid tissue. Hereditary motor and sensory neuropathy with proximal dominant involvement (HMSN-P) is an autosomal-dominant neurodegenerative disorder characterized by widespread fasciculations, proximal-predominant muscle weakness, and atrophy followed by distal sensory involvement. Recent genetic investigation indicates that formation of TFG-containing cytoplasmic inclusions and concomitant mislocalization of TAR DNA-binding protein 43 kDa (TDP-43) underlie motor neuron degeneration in HMSN-P. Pathological overlap of proteinopathies involving TFG and TDP-43 highlights a new pathway leading to motor neuron degeneration.		
Background Information	melanoma. Defects in TFG are a c typically arises as an irregular, so sensory neuropathy with proxima disorder characterized by widesp followed by distal sensory involv cytoplasmic inclusions and conce motor neuron degeneration in HM	lid or cystic mass from otherwise n l dominant involvement (HMSN-P) read fasciculations, proximal-predo ement. Recent genetic investigatio mitant mislocalization of TAR DNA ISN-P. Pathological overlap of prote	ormal thyroid tissue. Hereditary motor and is an autosomal-dominant neurodegenerativ minant muscle weakness, and atrophy n indicates that formation of TFG-containing -binding protein 43 kDa (TDP-43) underlie
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For technical support and original validation data for this product please contact: E: Proteintech-CN@ptglab.com T: 4006900926 W: ptgcn.com

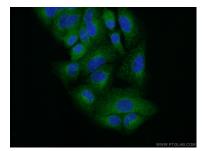
This product is exclusively available under Proteintech Group brand and is not available to purchase from any other manufacturer.

## Selected Validation Data





Immunohistochemical analysis of paraffinembedded human gliomas using 11571-1-AP (TFG antibody) at dilution of 1:100 (under 10x lens).



Immunofluorescent analysis of (10% Formaldehyde) fixed A549 cells using 11571-1-AP (TFG antibody) at dilution of 1:50 and Alexa Fluor 488-conjugated Goat Anti-Rabbit IgG(H+L).

