For Research Use Only

PARK7/DJ-1 Monoclonal antibody, PBS Only



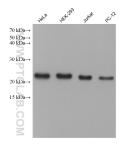
Catalog Number:68915-6-PBS

Basic Information	Catalog Number: 68915-6-PBS	GenBank Accession Number: BC008188	Purification Method: Protein A purification
	Concentration: 1 mg/ml	GeneID (NCBI): 11315	CloneNo.: 4G4E7
	Source: Mouse	UNIPROT ID: Q99497	
	Isotype: IgG1 Immunogen Catalog Number: AG28526	Full Name: Parkinson disease (autosomal recessive, early onset) 7 Calculated MW: 189 aa, 20 kDa	
		Applications	Tested Applications: WB, FC (Intra), Indirect ELISA
Species Specificity: human, mouse, rat, pig, rabbit			
Background Information	PARK7, also named as DJ1, belongs to the peptidase C56 family. It protects cells against oxidative stress and cell death. PARK7 plays a role in regulating expression or stability of the mitochondrial uncoupling proteins SLC25A14 and SLC25A27 in dopaminergic neurons of the substantia nigra pars compacta and attenuates the oxidative stress induced by calcium entry into the neurons via L-type channels during pacemaking. It eliminates hydrogen peroxide and protects cells against hydrogen peroxide-induced cell death. PARK7 has cell-growth promoting activity and transforming activity. It may function as a redox-sensitive chaperone. It's precursor undergoes a cleavage of a C-terminal peptide and subsequent activation of protease activity in response to oxidative stress. The amino acid replace at 166 (L \rightarrow P) reduces PARK7 protein stability and leads to increased degradation. The predicted MW of this protein is 20 kDa, An additional 25 kDa band can be observed due to modification (PMID: 31767755).		
Storage	Storage: Store at -80°C. The product is shipped with ice pa Storage Buffer: PBS Only	cks. Upon receipt, store it immediatel	y at −80°C

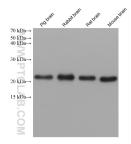
For technical support and original validation data for this product please contact:T: 4006900926E: Proteintech-CN@ptglab.comW: ptgcn.com

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

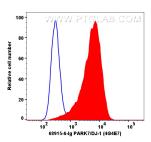
Selected Validation Data



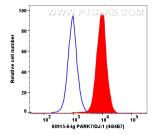
Various lysates were subjected to SDS PAGE followed by western blot with 68915-6-1g (PARK7/DJ-1 antibody) at dilution of 1:10000 incubated at room temperature for 1.5 hours. This data was developed using the same antibody clone with 68915-6-PBS in a different storage buffer formulation.



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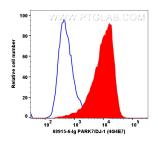


1x10^6 SKOV-3 cells were intracellularly stained with 0.2 µ g PARK7/DJ-1 Monoclonal antibody (68915-6-Ig, Clone: 4G4E7, red) and CoraLite® Plus 647-Goat Anti-Mouse Recombinant Secondary Antibody (H+L) (Cat.NO.RGAM005). Mouse IgG1 isotype control (66360-1-Ig, Clone: 1F8D3, blue) was parallel stained as control. Cells were fixed with 4% PFA. This data was developed using the same antibody clone with 68915-6-PBS in a different storage buffer formulation.



1x10^6 HeLa cells were intracellularly stained with 0.2 μ g PARK7/DJ-1 Monoclonal antibody (68915-6-Ig, Clone: 4G4E7, red) and CoraLite® Plus 647-Goat Anti-Mouse Recombinant Secondary Antibody (H+L) (Cat.NO.RGAM005). Mouse IgG1 isotype control (66360-1-Ig, Clone: 1F8D3, blue) was parallel stained as control. Cells were fixed with 4% PFA. This data was developed using the same antibody clone with 68915-6-PBS in a different storage buffer formulation.

1x10^6 Jurkat cells were intracellularly stained with 0.2 µ g PARK7/DJ-1 Monoclonal antibody (68915-6-Ig, Clone: 4G4E7, red) and CoraLite® Plus 647-Goat Anti-Mouse Recombinant Secondary Antibody (H+L) (Cat.NO.RGAM005). Mouse IgG1 isotype control (66360-1-Ig, Clone: 1F8D3, blue) was parallel stained as control. Cells were fixed with 4% PFA. This data was developed using the same antibody clone with 68915-6-PBS in a different storage buffer formulation.



1x10^6 Raji cells were intracellularly stained with 0.2 μ g PARK7/DJ-1 Monoclonal antibody (68915-6-Ig, Clone: 4G4E7, red) and CoraLite® Plus 647-Goat Anti-Mouse Recombinant Secondary Antibody (H+L) (Cat.NO.RGAM005). Mouse IgG1 isotype control (66360-1-Ig, Clone: 1F8D3, blue) was parallel stained as control. Cells were fixed with 4% PFA. This data was developed using the same antibody clone with 68915-6-PBS in a different storage buffer formulation.