bs-17657R

- DATASHEET ------

[Primary Antibody]

SPG20/Spartin Rabbit pAb



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Host: Rabbit	Isotype: IgG	Applications: IHC-P (1:100-500)
Clonality: Polyclonal		IHC-F (1:100-500) IF (1:100-500)
GenelD: 23111	SWISS: Q8N0X7	ICC/IF (1:100-500)
Target: SPG20/Spartin		Reactivity: (predicted: Human, Mouse, Rat, Rabbit)
Immunogen: KLH conjugated synthetic peptide derived from human SPG20/Spartin: 131-230/666.		
Purification: affinity purified by	Protein A	
Concentration: 1mg/ml		Predicted MW.: ^{73 kDa}
Storage: 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol. Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles.		Subcellular Cell membrane ,Cytoplasm Location: ,Nucleus
Background: This gene encodes a protein containing a MIT (Microtubule Interacting and Trafficking molecule) domain, and is implicated in regulating endosomal trafficking and mitochondria function. The protein localizes to mitochondria and partially co-localizes with microtubules. Stimulation with epidermal growth factor (EGF) results in protein translocation to the plasma membrane, and the protein functions in the degradation and intracellular trafficking of EGF receptor. Multiple alternatively spliced variants, encoding the same protein, have been identified. Mutations associated with this gene cause autosomal recessive spastic paraplegia 20 (Troyer syndrome). [provided by RefSeq, Nov 2008]		