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## ARHGAP36 Rabbit pAb

Catalog Number: bs-18116R
Target Protein: ARHGAP36

Concentration: 1mg/ml

Form: Liquid

Host: Rabbit
Clonality: Polyclonal

Isotype: IgG

Applications: IHC-P (1:100-500), IHC-F (1:100-500), IF (1:100-500), ICC/IF (1:100-500), ELISA (1:5000-10000)

Reactivity: (predicted:Human, Mouse, Rat)

Predicted MW: 57 kDa Entrez Gene: 158763 Swiss Prot: Q6ZRI8

Source: KLH conjugated synthetic peptide derived from human ARHGAP36: 351-450/547.

Purification: affinity purified by Protein A

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.

Background: ARHGAP36 is a 547 amino acid protein that contains one Rho-GAP domain. Conserved in

 $chimpanzee, dog, cow, mouse \ and \ rat, ARHGAP36 \ exists \ as \ five \ alternatively \ spliced$ 

isoforms and is encoded by a gene that maps to human chromosome Xq26.1. Chromosome X consists of nearly 153 million base pairs encoding approximately 1,000 genes. More than one copy of the X chromosome with a Y chromosome causes Klinefelter's syndrome. A single copy of X alone leads to Turner's syndrome. More than 2 copies of the X chromosome, in the absence of a Y chromosome, is known as Triple X syndrome. Color blindness, hemophilia, and Duchenne muscular dystrophy are X chromosome-linked conditions that affect males

more frequently because males carry a single X chromosome.