

Myosin IIa (phospho-Ser1943) rabbit pAb

Catalog_no :	AP1409
Applications :	WB
Reactivity :	Human,Mouse,Rat
Category :	抗原抗体
Size :	100µg/50µg/20µg
Gene_name :	MYH9
Protein_name :	Myosin IIa (Ser1943)
Humangene_id :	4627
Humanswissprot_no :	P35579
Mousegene_id :	17886
Mouseswissprot_no :	Q8VDD5
Ratgene_id :	25745
Ratswissprot_no :	Q62812
Immunogen :	Synthesized phosho peptide around human Myosin IIa (Ser1943)
Specificity :	This antibody detects endogenous levels of Human Mouse Rat Myosin IIa (phospho-Ser1943)
Formulation :	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
Source :	Rabbit
Dilution :	WB 1:1000-2000
Purification :	The antibody was affinity-purified from rabbit serum by affinity-chromatography using specific immunogen.
Concentration :	1 mg/ml
Storage_stability :	-20°C/1 year
Other_name :	Myosin-9 (Cellular myosin heavy chain, type A) (Myosin heavy chain 9) (Myosin heavy chain, non-muscle IIa) (Non-muscle myosin heavy chain A) (NMMHC-A) (Non-muscle

myosin heavy chain IIa) (NMMHC II-a) (NMMHC-IIA)

Molecular Weight : 215KD

Background : myosin heavy chain 9(MYH9) Homo sapiens This gene encodes a conventional non-muscle myosin; this protein should not be confused with the unconventional myosin-9a or 9b (MYO9A or MYO9B). The encoded protein is a myosin IIA heavy chain that contains an IQ domain and a myosin head-like domain which is involved in several important functions, including cytokinesis, cell motility and maintenance of cell shape. Defects in this gene have been associated with non-syndromic sensorineural deafness autosomal dominant type 17, Epstein syndrome, Alport syndrome with macrothrombocytopenia, Sebastian syndrome, Fechtner syndrome and macrothrombocytopenia with progressive sensorineural deafness. [provided by RefSeq, Dec 2011],
