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## MYH9 Antibody

cycles.Purification MethodAntigen Affinity PurifiedIsotypeIgGAliasmyosin, heavy chain 9, non- muscle;MYH9;DFNA17;EPSTS;FTNS;MGC104539;MHA;NN ;Product TypePurified Rabbit Anti human PolyClonal AntibodyImmunogen SpeciesHomo sapiens (Human)Target NamesMYH9Target DetailsThis gene encodes a myosin IIA heavy chain that contains an myosin head-like domain. The protein is involved in several in including cytokinesis, cell motility and maintenance of cell sh MYH9 are the cause of non-syndrome, Alport syndrome with		
Uniprot No.P35579ImmunogenHuman MYH9Raised InRabbitSpecies ReactivityHuman,Mouse,RatTested ApplicationsELISA,WB,IHC,IFStorage BufferPBS with 0.1% Sodium Azide, 50% Glycerol, pH 7.320°C, cycles.Purification MethodAntigen Affinity PurifiedIsotypeIgGAliasmyosin, heavy chain 9, non- muscle;MYH9;DFNA17;EPSTS;FTNS;MGC104539;MHA;NM ;Product TypePurified Rabbit Anti human PolyClonal AntibodyImmunogen SpeciesHomo sapiens (Human)Target NamesMYH9Target DetailsThis gene encodes a myosin IIA heavy chain that contains an myosin head-like domain. The protein is involved in several i including cytokinesis, cell motility and maintenance of cell sh MYH9 are the cause of non-syndromic sensorineural deafner dominant type 17, Epstein syndrome, Alport syndrome with	uct Code CS	SB-PA015303GA01HU
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	my inc M\ do ma	his gene encodes a myosin IIA heavy chain that contains an IQ domain and a yosin head-like domain. The protein is involved in several important functions, cluding cytokinesis, cell motility and maintenance of cell shape. Defects in YH9 are the cause of non-syndromic sensorineural deafness autosomal pminant type 17, Epstein syndrome, Alport syndrome with acrothrombocytopenia, Sebastian syndrome, Fechtner syndrome and acrothrombocytopenia with progressive sensorineural deafness.

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