

## ARG56246 anti-MYH9 antibody

Package: 100 µl  
Store at: -20°C

### Summary

Product Description	Rabbit Polyclonal antibody recognizes MYH9
Tested Reactivity	Hu, Ms
Tested Application	IHC-P, WB
Host	Rabbit
Clonality	Polyclonal
Isotype	IgG
Target Name	MYH9
Species	Human
Immunogen	Recombinant protein of Human MYH9
Conjugation	Un-conjugated
Alternate Names	NMHC-II-A; Myosin heavy chain 9; FTNS; Non-muscle myosin heavy chain IIa; Non-muscle myosin heavy chain A; BDPLT6; EPSTS; NMMHCA; MHA; Myosin-9; DFNA17; Myosin heavy chain, non-muscle IIa; NMMHC-IIA; NMMHC II-a; NMMHC-A; Cellular myosin heavy chain, type A

### Application Instructions

Application table	Application	Dilution
	IHC-P	1:50 - 1:100
	WB	1:500 - 1:1000
Application Note	* The dilutions indicate recommended starting dilutions and the optimal dilutions or concentrations should be determined by the scientist.	
Positive Control	Jurkat	

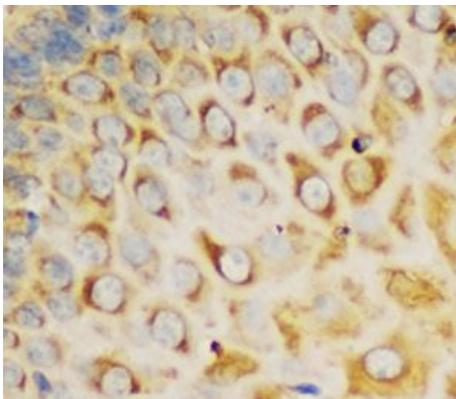
### Properties

Form	Liquid
Purification	Affinity purification with immunogen.
Buffer	PBS (pH 7.3), 0.02% Sodium azide and 50% Glycerol.
Preservative	0.02% Sodium azide
Stabilizer	50% Glycerol
Storage instruction	For continuous use, store undiluted antibody at 2-8°C for up to a week. For long-term storage, aliquot and store at -20°C. Storage in frost free freezers is not recommended. Avoid repeated freeze/thaw cycles. Suggest spin the vial prior to opening. The antibody solution should be gently mixed before use.
Note	For laboratory research only, not for drug, diagnostic or other use.

## Bioinformation

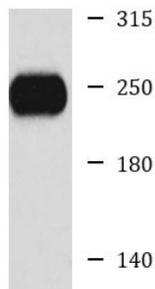
Database links	<a href="#">GeneID: 17886 Mouse</a> <a href="#">GeneID: 4627 Human</a> <a href="#">Swiss-port # P35579 Human</a> <a href="#">Swiss-port # Q8VDD5 Mouse</a>
Gene Symbol	MYH9
Gene Full Name	myosin, heavy chain 9, non-muscle
Background	This gene encodes a conventional non-muscle myosin; this 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]
Function	Cellular myosin that appears to play a role in cytokinesis, cell shape, and specialized functions such as secretion and capping. During cell spreading, plays an important role in cytoskeleton reorganization, focal contacts formation (in the margins but not the central part of spreading cells), and lamellipodial retraction; this function is mechanically antagonized by MYH10. [UniProt]
Calculated Mw	227 kDa
PTM	ISGylated.

## Images



ARG56246 anti-MYH9 antibody IHC-P image

Immunohistochemistry: Paraffin-embedded Human esophagus stained with ARG56246 anti-MYH9 antibody at 1:100 dilution.



Jurkat

ARG56246 anti-MYH9 antibody WB image

Western blot: Jurkat cell lysate stained with ARG56246 anti-MYH9 antibody.