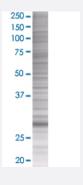


MYO3A 293T Cell Transient Overexpression Lysate(Denatured)

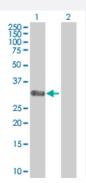
Catalog # H00053904-T01 Size 100 uL

Applications



SDS-PAGE Gel

MYO3A transfected lysate.



Western Blot

Lane 1: MYO3A transfected lysate (27.6 KDa)

Lane 2: Non-transfected lysate.

Specification	
Transfected Cell Line	293T
Plasmid	pCMV-MYO3A full-length
Host	Human
Theoretical MW (kDa)	27.28
Interspecies Antigen Sequence	Mouse (94)



Product Information

Quality Control Testing	Transient overexpression cell lysate was tested with Anti-MYO3A antibody (H00053904-B01) by We stern Blots. SDS-PAGE Gel MYO3A transfected lysate. Western Blot Lane 1: MYO3A transfected lysate (27.6 KDa) Lane 2: Non-transfected lysate.
Storage Buffer	1X Sample Buffer (50 mM Tris-HCl, 2% SDS, 10% glycerol, 300 mM 2-mercaptoethanol, 0.01% Bro mophenol blue)
Storage Instruction	Store at -80°C. Aliquot to avoid repeated freezing and thawing.

Applications

Western Blot

Gene Info — MYO3A	
Entrez GenelD	<u>53904</u>
GeneBank Accession#	BC045538.1
Protein Accession#	=
Gene Name	MYO3A
Gene Alias	DFNB30
Gene Description	myosin IIIA
Omim ID	<u>606808</u> <u>607101</u>
Gene Ontology	<u>Hyperlink</u>
Gene Summary	The protein encoded by this gene belongs to the myosin superfamily. Myosins are actin-depende nt motor proteins and are categorized into conventional myosins (class II) and unconventional myosins (classes I and III through XV) based on their variable C-terminal cargo-binding domains. Class III myosins, such as this one, have a kinase domain N-terminal to the conserved N-terminal motor domains and are expressed in photoreceptors. The protein encoded by this gene plays an important role in hearing in humans. Three different recessive, loss of function mutations in the encoded protein have been shown to cause nonsyndromic progressive hearing loss. Expression of this gene is highly restricted, with the strongest expression in retina and cochlea. [provided by RefSeq
Other Designations	OTTHUMP00000019339



Disease

- Alzheimer Disease
- Anxiety Disorders
- Arthritis
- Colorectal Neoplasms
- Depressive Disorder
- Genetic Predisposition to Disease
- Kidney Failure