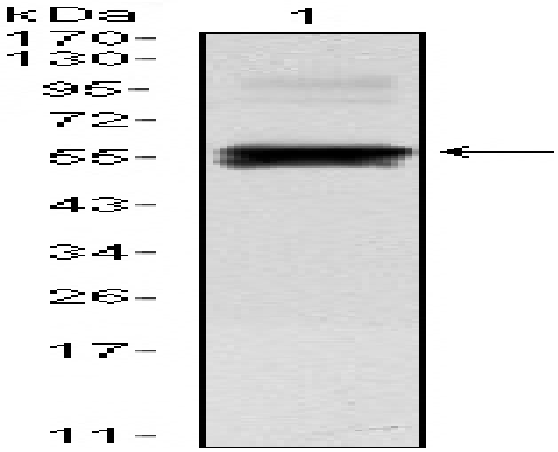
Best Tools for immunology Research

Pax-6 Monoclonal Antibody

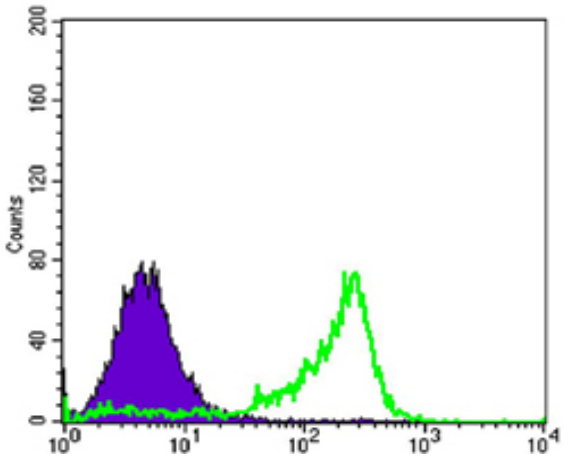
Catalog No :	YM0508
Reactivity :	Human
Applications :	WB;FCM;ELISA
Target :	Pax-6
Fields :	>>Signaling pathways regulating pluripotency of stem cells;>>Maturity onset diabetes of the young
Gene Name :	PAX6
Protein Name :	Paired box protein Pax-6
Human Gene Id :	5080
Human Swiss Prot No :	P26367
Mouse Swiss Prot No :	P63015
Immunogen :	Purified recombinant fragment of human Pax-6 expressed in E. Coli.
Specificity :	Pax-6 Monoclonal Antibody detects endogenous levels of Pax-6 protein.
Formulation :	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
Source :	Monoclonal, Mouse
Dilution :	WB 1:500 - 1:2000. Flow cytometry: 1:200 - 1:400. ELISA: 1:10000. Not yet tested in other applications.
Purification :	Affinity purification
Storage Stability :	-15°C to -25°C/1 year(Do not lower than -25°C)
Molecularweight :	47kD

Cell Pathway :	Maturity onset diabetes of the young;
P References :	1.Invest Ophthalmol Vis Sci. 2009 Jun;50(6):2581-90. 2.J Biol Chem. 2009 Oct 2;284(40):27524-32. 3.J Biol Chem. 2010 Jan 22;285(4):2527-36.
Background :	This gene encodes a homeobox and paired domain-containing protein that binds DNA and functions as a regulator of transcription. Activity of this protein is key in the development of neural tissues, particularly the eye. This gene is regulated by multiple enhancers located up to hundreds of kilobases distant from this locus. Mutations in this gene or in the enhancer regions can cause ocular disorders such as aniridia and Peter's anomaly. Use of alternate promoters and alternative splicing result in multiple transcript variants encoding different isoforms. [provided by RefSeq, Jul 2015],
Function :	developmental stage:Expressed in the developing eye and brain.,disease:Defects in PAX6 are a cause of autosomal dominant keratitis [MIM:148190]. It is an eye disorder characterized by corneal opacification and vascularization, and by foveal hypoplasia.,disease:Defects in PAX6 are a cause of bilateral optic nerve hypoplasia [MIM:165550]; also known as bilateral optic nerve aplasia. Inheritance is autosomal dominant.,disease:Defects in PAX6 are a cause of coloboma of optic nerve [MIM:120430].,disease:Defects in PAX6 are a cause of ectopia pupillae [MIM:129750]. It is a congenital eye malformation in which the pupils are displaced from their normal central position.,disease:Defects in PAX6 are a cause of foveal hypoplasia [MIM:136520]. Foveal hypoplasia can be isolated or associated with presenile cataract. Inheritance is autosomal dominant.,disease:Defects in PAX6 are a cause of Gillespie
Subcellular Location :	Nucleus .; [Isoform 1]: Nucleus .; [Isoform 5a]: Nucleus .
Expression :	[Isoform 1]: Expressed in lymphoblasts. ; [Isoform 5a]: Weakly expressed in lymphoblasts.
Sort :	11652
No4 :	1
Host :	Mouse
Modifications :	Unmodified

Products Images



Western Blot analysis using Pax-6 Monoclonal Antibody against recombinant Pax-6 protein (1).



Flow cytometric analysis of 3T3-L1 cells using Pax-6 Monoclonal Antibody (green) and negative control (purple).