



RAX2 mouse mAb

Catalog No	YP-mAb-11192
Isotype	IgG
Reactivity	Human;Mouse;Rat
Applications	WB
Gene Name	RAX2 QRX RAXL1
Immunogen	Synthesized peptide derived from human RAX2 AA range: 95-145
Specificity	This antibody detects endogenous levels of RAX2 at Human
Formulation	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
Source	Monoclonal, Mouse,IgG
Purification	The antibody was affinity-purified from mouse antiserum by affinity-chromatography using epitope-specific immunogen.
Dilution	WB 1:500-1:2000
Concentration	1 mg/ml
Purity	≥90%
Storage Stability	-20°C/1 year
Synonyms	ARMD6;CORD11;QRX;RAX2;RAXL1;RAX2 QRX RAXL1
Observed Band	
Calculated Molecular Weight	
Cell Pathway	Nucleus .
Tissue Specificity	
Function	disease:Defects in RAX2 are the cause of age-related macular degeneration type 6 (ARMD6) [MIM:603075]. ARMD is in most patients manifest as ophthalmoscopically visible yellowish accumulations of protein and lipid (known as drusen) that lie beneath the retinal pigment epithelium and within an elastin-containing structure known as Bruch's membrane. ARMD is likely to be a mechanistically heterogeneous group of disorders.,disease:Defects in RAX2 are the cause of cone-rod dystrophy type 11 (CORD11) [MIM:610381]. CORD is characterized by the initial degeneration of cone photoreceptor cells, thus causing early loss of visual acuity and color vision, followed by the degeneration of rod photoreceptor cells and leading to progressive night blindness and peripheral visual field loss.,domain:The Homeobox transactivates the Ret-1 element in the presence of CRX and NRL.,function:May be involved in mod



Background

This gene encodes a homeodomain-containing protein that plays a role in eye development. Mutation of this gene causes age-related macular degeneration type 6, an eye disorder resulting in accumulations of protein and lipid beneath the retinal pigment epithelium and within the Bruch's membrane. Defects in this gene can also cause cone-rod dystrophy type 11, a disease characterized by the initial degeneration of cone photoreceptor cells and resulting in loss of color vision and visual acuity, followed by the degeneration of rod photoreceptor cells, which progresses to night blindness and the loss of peripheral vision. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Jan 2016],

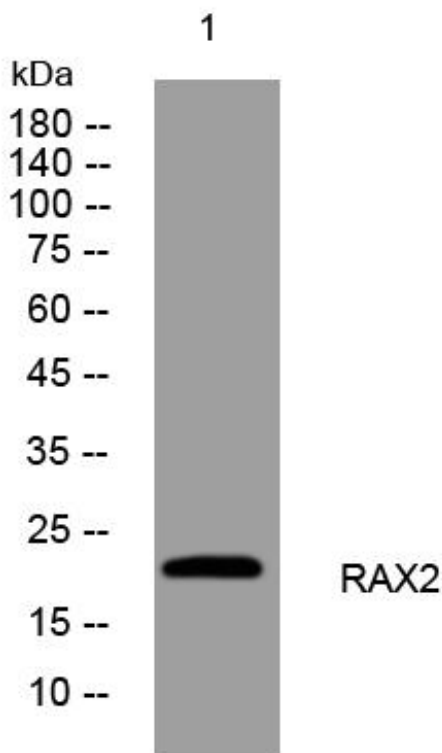
matters needing attention

Avoid repeated freezing and thawing!

Usage suggestions

This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.

Products Images



Western Blot analysis of various cells using RAX2 mouse mAb