

## FoxC1/2 rabbit pAb

Cat No.: ES5279

For research use only

## Overview

**Product Name** FoxC1/2 rabbit pAb

**Host species** Rabbit

WB;IHC;IF;ELISA **Applications Species Cross-Reactivity** Human; Mouse; Rat

**Recommended dilutions** Western Blot: 1/500 - 1/2000.

Immunohistochemistry: 1/100 - 1/300.

Immunofluorescence: 1/200 - 1/1000. ELISA: 1/40000. Not yet tested in other applications. The antiserum was produced against synthesized

**Immunogen** peptide derived from human FOXC1/2. AA

range:151-200

**Specificity** FoxC1/2 Polyclonal Antibody detects endogenous

levels of FoxC1/2 protein.

**Formulation** Liquid in PBS containing 50% glycerol, 0.5% BSA and

0.02% sodium azide.

Store at -20°C. Avoid repeated freeze-thaw cycles. **Storage** 

**Protein Name** Forkhead box protein C1/2

**Gene Name** FOXC1/FOXC2

Cellular localization Nucleus . Colocalizes with PITX2 isoform 3 in the

> nucleus at subnuclear chromatine regions (PubMed:16449236). Colocalizes with CBX5 to a heterochromatin-rich region of the nucleus (PubMed:15684392). Colocalizes with GLI2 in the

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nucleus (By similarity). .

Purification The antibody was affinity-purified from rabbit

antiserum by affinity-chromatography using

epitope-specific immunogen.

Clonality Polyclonal Concentration 1 mg/ml **Observed band** 57kD **Human Gene ID** 2296/2303

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**Human Swiss-Prot Number** Q12948/Q99958

**Alternative Names** FOXC1; FKHL7; FREAC3; Forkhead box protein C1;

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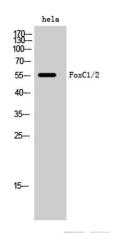


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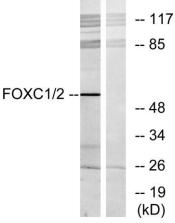


**Background** 

Forkhead-related protein FKHL7; Forkhead-related transcription factor 3; FREAC-3; FOXC2; FKHL14; MFH1; Forkhead box protein C2; Forkhead-related protein FKHL14; Mesenchyme fork head protein 1; This gene belongs to the forkhead family of transcription factors which is characterized by a distinct DNA-binding forkhead domain. The specific function of this gene has not yet been determined; however, it has been shown to play a role in the regulation of embryonic and ocular development. Mutations in this gene cause various glaucoma phenotypes including primary congenital glaucoma, autosomal dominant iridogoniodysgenesis anomaly, and Axenfeld-Rieger anomaly. [provided by RefSeq, Jul 2008],



Western Blot analysis of hela cells using FoxC1/2 Polyclonal Antibody diluted at 1:2000 cells nucleus extracted by Minute TM Cytoplasmic and Nuclear Fractionation kit (SC-003,Inventbiotech,MN,USA).



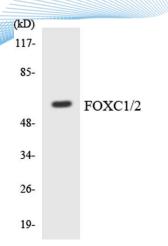
Western blot analysis of lysates from RAW264.7 cells, using FOXC1/2 Antibody. The lane on the right is blocked with the synthesized peptide.



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Western blot analysis of the lysates from Jurkat cells using FOXC1/2 antibody.



Immunohistochemical analysis of paraffin-embedded human tonsil. 1, Tris-EDTA,pH9.0 was used for antigen retrieval. 2 Antibody was diluted at 1:200(4° overnight.3,Secondary antibody was diluted at 1:200(room temperature, 45min).

