

For Research Use Only

# FOXC2 Polyclonal antibody

Catalog Number: 23066-1-AP

Featured Product

6 Publications



## Basic Information

### Catalog Number:

23066-1-AP

### Size:

150ul, Concentration: 300 µg/ml by Nanodrop and 293 µg/ml by Bradford method using BSA as the standard;

### Source:

Rabbit

### Isotype:

IgG

### Immunogen Catalog Number:

AG19378

### GenBank Accession Number:

BC113437

### GeneID (NCBI):

2303

### Full Name:

forkhead box C2 (MFH-1, mesenchyme forkhead 1)

### Calculated MW:

501 aa, 54 kDa

### Observed MW:

56 kDa

### Purification Method:

Antigen affinity purification

### Recommended Dilutions:

WB 1:500-1:2000

IF 1:50-1:500

## Applications

### Tested Applications:

IF, WB, ELISA

### Cited Applications:

IF, IHC, WB

### Species Specificity:

human, rat, mouse

### Cited Species:

human, mouse, rat

### Positive Controls:

WB: A375 cells, A431 cells, rat spleen tissue

IF: A375 cells,

## Background Information

Transcriptional activator. Might be involved in the formation of special mesenchymal tissues. Defects in FOXC2 are the cause of lymphedema hereditary type 2 (LMPH2) [MIM:153200]; also known as Meige lymphedema. Hereditary lymphedema is a chronic disabling condition which results in swelling of the extremities due to altered lymphatic flow. Patients with lymphedema suffer from recurrent local infections, and physical impairment. Defects in FOXC2 are a cause of lymphedema-yellow nails (LYYN) [MIM:153300]. LYYN is characterized by yellow, dystrophic, thick and slowly growing nails, associated with lymphedema and respiratory involvement. Lymphedema occurs more often in the lower limbs. It can appear at birth or later in life. Onset generally follows the onset of ungual abnormalities. Defects in FOXC2 are a cause of lymphedema-distichiasis (LYD) [MIM:153400]. LYD is characterized by primary limb lymphedema usually starting at puberty (but in some cases later or at birth) and associated with distichiasis (double rows of eyelashes, with extra eyelashes growing from the Meibomian gland orifices). This antibody specifically recognizes the 56 kDa FOXC2 protein.

## Notable Publications

Author	Pubmed ID	Journal	Application
Fei-Xiang Lin	27904711	Am J Transl Res	WB
Mingliang Liu	31367258	Theranostics	WB
Ping Wang	25662617	Acta Biochim Biophys Sin (Shanghai)	WB

## Storage

### Storage:

Store at -20°C. Stable for one year after shipment.

### Storage Buffer:

PBS with 0.02% sodium azide and 50% glycerol pH 7.3.

Aliquoting is unnecessary for -20°C storage

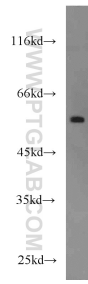
For technical support and original validation data for this product please contact:

T: 1 (888) 4PTGLAB (1-888-478-4522) (toll free in USA), or 1(312) 455-8498 (outside USA)

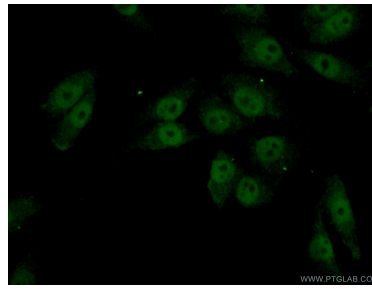
E: [proteintech@ptglab.com](mailto:proteintech@ptglab.com)  
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## Selected Validation Data



A375 cells were subjected to SDS PAGE followed by western blot with 23066-1-AP (FOXC2 antibody) at dilution of 1:1000 incubated at room temperature for 1.5 hours.



Immunofluorescent analysis of (4% PFA) fixed A375 cells using 23066-1-AP (FOXC2 antibody) at dilution of 1:50 and Alexa Fluor 488-conjugated AffiniPure Goat Anti-Rabbit IgG(H+L).