For Research Use Only

FOXC1 Monoclonal antibody

Catalog Number:66568-1-lg Featured Product



Basic Information

Catalog Number: 66568-1-Ig

Size: 1500 ug/ml

Source:

Mouse

Isotype:

NM_001453 GeneID (NCBI): 2296 UNIPROT ID:

GenBank Accession Number:

Q12948
Full Name:
forkhead box C1
Calculated MW:

57 kDa Observed MW: 70-75 kDa Purification Method:

Protein G purification CloneNo.:

1F4E11 Recommended Dilutions: WB 1:2000-1:20000

Applications

Tested Applications:

WB,ELISA

Species Specificity:

human

Positive Controls:

WB: HeLa cells, HEK-293 cells, HEK293 cells, LO2 cells,

HepG2 cells

Background Information

FOXC1, also named as FKHL7 and FREAC3, binding of FREAC-3 and FREAC-4 to their cognate sites results in bending of the DNA at an angle of 80-90 degrees. Defects in FOXC1 are the cause of Axenfeld-Rieger syndrome type 3 (RIEG3). Defects in FOXC1 are the cause of iridogoniodysgenesis anomaly (IGDA). Defects in FOXC1 are a cause of Peters anomaly.

Storage

Storage:

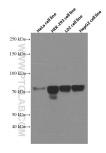
Store at -20°C. Stable for one year after shipment.

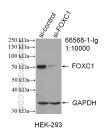
Storage Buffer:

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

Aliquoting is unnecessary for -20°C storage

Selected Validation Data





Various cells were subjected to SDS PAGE followed by western blot with 66568-1-1g (FOXC1 antibody) at dilution of 1:10000 incubated at room temperature for 1.5 hours.

WB result of FOXC1 antibody (66568-1-lg; 1:10000; incubated at room temperature for 1.5 hours) with sh-Control and sh-FOXC1 transfected HEK-293 cells.