

# Anti-FOG2 Antibody

Rabbit polyclonal antibody to FOG2 Catalog # AP61530

#### **Product Information**

ApplicationWB, IHCPrimary AccessionQ8WW38ReactivityHumanHostRabbitClonalityPolyclonalCalculated MW128159

#### **Additional Information**

**Gene ID** 23414

Other Names FOG2; ZNF89B; Zinc finger protein ZFPM2; Friend of GATA protein 2; FOG-2;

Friend of GATA 2; hFOG-2; Zinc finger protein 89B; Zinc finger protein

multitype 2

**Target/Specificity** KLH-conjugated synthetic peptide encompassing a sequence within the center

region of human FOG2. The exact sequence is proprietary.

**Dilution** WB~~WB (1/500 - 1/1000), IHC (1/100 - 1/200) IHC~~WB (1/500 - 1/1000), IHC

(1/100 - 1/200)

**Format** Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30%

glycerol, and 0.09% (W/V) sodium azide.

**Storage** Store at -20 °C.Stable for 12 months from date of receipt

#### **Protein Information**

Name ZFPM2

**Synonyms** FOG2, ZNF89B

**Function** Transcription regulator that plays a central role in heart morphogenesis and

development of coronary vessels from epicardium, by regulating genes that are essential during cardiogenesis. Essential cofactor that acts via the formation of a heterodimer with transcription factors of the GATA family GATA4, GATA5 and GATA6. Such heterodimer can both activate or repress transcriptional activity, depending on the cell and promoter context. Also required in gonadal differentiation, possibly be regulating expression of SRY.

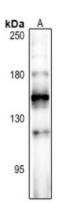
Probably acts a corepressor of NR2F2 (By similarity).

Cellular Location Nucleus.

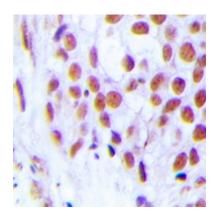
## **Background**

KLH-conjugated synthetic peptide encompassing a sequence within the center region of human FOG2. The exact sequence is proprietary.

### **Images**



Western blot analysis of FOG2 expression in Beas2B (A) whole cell lysates.



Immunohistochemical analysis of FOG2 staining in human breast cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.