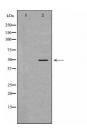


Affinity Biosciences website:www.affbiotech.com order:order@affbiotech.com

GATA2 Ab

| Cat.#: DF6117<br>Size: 100ul,200ul | Concn.: 1mg/ml<br>Source: Rabbit  | Mol.Wt.: 49kDa<br>Clonality: Polyclonal |
|------------------------------------|---|---|
| Application:                       | WB 1:500-1:2000 IHC 1:50-1:200, IF/ICC 1:100-1:500  |   |
| Reactivity:                        | Human,Mouse,Rat   |   |
| Purification:                      | The antiserum was purified by p<br>chromatography using SulfoLinl<br>Fisher Scientific).  | , ,                                     |
| Specificity:                       | GATA2 Ab detects endogenous levels of total GATA2.  |   |
| Immunogen:                         | A synthesized peptide derived from human GATA2.   |   |
| Uniprot:                           | P23769  |   |
| Description:                       | The zinc finger transcription factor GATA-2 is widely<br>expressed and plays an essential role in many<br>developmental processes (1). Studies on GATA-2 knockout<br>mice indicate that this protein is required in hematopoiesis<br>(2). GATA-2 also inhibits the differentiation of white (3) and<br>brown adipocytes (4) and has been shown to suppress the<br>proliferation of neuronal progenitor cells (5). |   |
| Subcellular Location:              | Nucleus.  |   |
| Tissue Specificity:                | Endothelial cells.  |   |
| Storage Condition and<br>Buffer:   | Rabbit IgG in phosphate buffered saline , pH 7.4, 150mM<br>NaCl, 0.02% sodium azide and 50% glycerol.Store at -20<br>°C.Stable for 12 months from date of receipt.  |   |



Western blot analysis of Jurkat lysates using GATA2 Ab. The lane on the left was treated with the antigen-specific peptide.





DF6117 at 1/100 staining Mouse liver tissue by IHC-P. The sample was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The sample was then blocked and incubated with the Ab for 1.5 hours at 22°C. An HRP conjugated goat anti-rabbit Ab was used as the secondary.



DF6117 staining HepG2 by IF/ICC. The sample were fixed with PFA and permeabilized in 0.1% Triton X-100,then blocked in 10% serum for 45 minutes at 25°C. The primary Ab was diluted at 1/200 and incubated with the sample for 1 hour at 37°C. An Alexa Fluor 594 conjugated goat anti-rabbit IgG (H+L) Ab, diluted at 1/600, was used as the secondary Ab.

<code>IMPORTANT:</code> For western blot, incubate membrane with diluted primary Ab in 5% w/v milk , 1X TBS, 0.1% Tween®20 at 4°C with gentle shaking, overnight.

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