

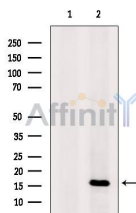
## GYPA Ab

Cat.#: DF10170  
Size: 100ul,200ul

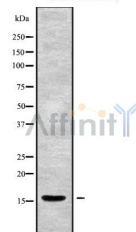
Concn.: 1mg/ml  
Source: Rabbit

Mol.Wt.: 16 kDa  
Clonality: Polyclonal

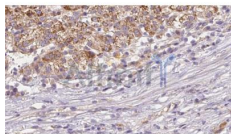
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|-------------------------------|--|
| Application:                  | WB 1:1000-3000 IHC 1:200, IF/ICC 1:100-1:500   |
| Reactivity:                   | Human,Rat  |
| Purification:                 | The antiserum was purified by peptide affinity chromatography using SulfoLink™ Coupling Resin (Thermo Fisher Scientific).                                    |
| Specificity:                  | GYPA Ab detects endogenous levels of total GYPA.   |
| Immunogen:                    | A synthesized peptide derived from human GYPA.   |
| Uniprot:                      | P02724   |
| Subcellular Location:         | Cell membrane. Appears to be colocalized with SLC4A1.  |
| Similarity:                   | Belongs to the glycophorin A family.   |
| Storage Condition and Buffer: | Rabbit IgG in phosphate buffered saline , pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol.Store at -20 °C.Stable for 12 months from date of receipt. |



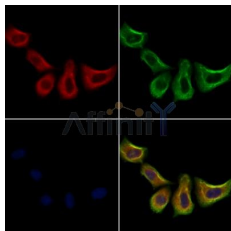
Western blot analysis of extracts from Rat muscle, using GYPA Ab. The lane on the left was treated with blocking peptide.



Western blot analysis GYPA using 293 whole cell lysates



DF10170 at 1/100 staining Human liver cancer 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.



DF10170 staining HepG2 cells by IF/ICC. The samples were fixed with PFA and permeabilized in 0.1% Triton X-100, then blocked in 10% serum for 45 minutes at 25°C. Samples were then incubated with primary Ab(DF10170 1:200) and mouse anti-beta tubulin Ab(T0023 1:200) for 1 hour at 37°C. An AlexaFluor594 conjugated goat anti-rabbit IgG(H+L) Ab(Red) and an AlexaFluor488 conjugated goat anti-mouse IgG(H+L) Ab(Green) were used as the secondary Ab.

**IMPORTANT:** 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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