

Datasheet for ABIN6257190

anti-FGF13 antibody (Internal Region)

1 Image



Go to Product page

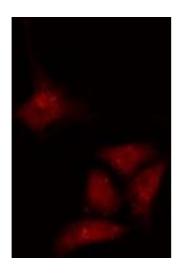
| Overview | |
|-----------------------|--|
| Quantity: | 100 μL |
| Target: | FGF13 |
| Binding Specificity: | Internal Region |
| Reactivity: | Human, Rat, Mouse |
| Host: | Rabbit |
| Clonality: | Polyclonal |
| Conjugate: | This FGF13 antibody is un-conjugated |
| Application: | Western Blotting (WB), ELISA, Immunocytochemistry (ICC), Immunofluorescence (IF) |
| Product Details | |
| lmmunogen: | A synthesized peptide derived from human FGF13, corresponding to a region within the internal amino acids. |
| Isotype: | IgG |
| Specificity: | FGF13 Antibody detects endogenous levels of total FGF13. |
| Predicted Reactivity: | Pig,Bovine,Horse,Sheep,Rabbit,Dog,Chicken,Xenopus |
| Purification: | The antiserum was purified by peptide affinity chromatography using SulfoLink TM Coupling Resin (Thermo Fisher Scientific). |
| Target Details | |
| Target: | FGF13 |

Target Details

Expiry Date:

| FGF13 (FGF13 Products) |
|--|
| Description: Microtubule-binding protein which directly binds tubulin and is involved in both polymerization and stabilization of microtubules. Through its action on microtubules, may participate to the refinement of axons by negatively regulating axonal and leading processes branching. Plays a crucial role in neuron polarization and migration in the cerebral cortex and the hippocampus. Gene: FGF13 |
| 28 kDa |
| 2258 |
| Q92913 |
| Regulation of Cell Size |
| |
| WB 1:500-1:1000, IF/ICC 1:100-1:500, ELISA(peptide) 1:20000-1:40000 |
| For Research Use only |
| |
| Liquid |
| 1 mg/mL |
| Rabbit IgG in phosphate buffered saline , pH 7.4, 150 mM NaCl, 0.02 % sodium azide and 50 % glycerol. |
| Sodium azide |
| This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only. |
| -20 °C |
| Store at -20 °C. Stable for 12 months from date of receipt. |
| |

12 months



Immunofluorescence (fixed cells)

Image 1. ABIN6275763 staining Hela cells 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¡ãC. The primary antibody was diluted at 1/200 and incubated with the sample for 1 hour at 37¡ãC. An Alexa Fluor 594 conjugated goat anti-rabbit IgG (H+L) antibody(Cat.# S0006), diluted at 1/600, was used as secondary antibod