## **SUZ12 Antibody**



## PACO46538

## **Product Information**

**Recommended dilutions:** 

Size: **Protein Background:** 

50ug Polycomb group (PcG) protein. Component of the PRC2/EED-EZH2 complex, which

methylates 'Lys-9' (H3K9me) and 'Lys-27' (H3K27me) of histone H3, leading to Reactivity: transcriptional repression of the affected target gene. The PRC2/EED-EZH2 complex

may also serve as a recruiting platform for DNA methyltransferases, thereby linking two Human, Rat

epigenetic repression systems. Genes repressed by the PRC2/EED-EZH2 complex

Source: include HOXC8, HOXA9, MYT1 and CDKN2A.

Rabbit Gene ID:

SUZ12 Isotype:

lgG Uniprot

Q15022 **Applications:** 

ELISA, WB, IHC, IF Synonyms:

Polycomb protein SUZ12 (Chromatin precipitated E2F target 9 protein) (ChET 9 protein)

(Joined to JAZF1 protein) (Suppressor of zeste 12 protein homolog), SUZ12, CHET9 ELISA:1:2000-1:10000, WB:1:500-1:5000, JJAZ1 KIAA0160

IHC:1:1000-1:2000, IF:1:50-1:500

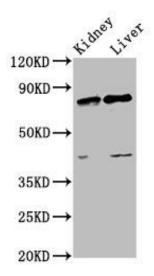
Immunogen:

Recombinant Human Polycomb protein SUZ12 protein (131-305AA).

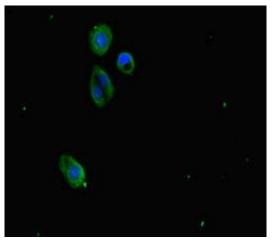
Storage:

Preservative: 0.03% Proclin 300. Constituents: 50% Glycerol, 0.01M PBS, PH 7.4

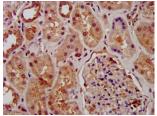
## **Product Images**



Western Blot. Positive WB detected in: Rat kidney tissue, Rat liver tissue. All lanes: SUZ12 antibody at 3.4µg/ml. Secondary. Goat polyclonal to rabbit IgG at 1/50000 dilution. Predicted band size: 84 kDa. Observed band size: 84 kDa.



Immunofluorescence staining of HepG2 cells with PACO46538 at 1:333, counter-stained with DAPI. The cells were fixed in 4% formaldehyde, permeabilized using 0.2% Triton X-100 and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4°C. The secondary antibody was Alexa Fluor 488-congugated AffiniPure Goat Anti-Rabbit IgG(H+L).



IHC image of PACO46538 diluted at 1:1000 and staining in paraffinembedded human kidney tissue performed on a Leica BondTM system. After dewaxing and hydration, antigen retrieval was mediated by high pressure in a citrate buffer (pH 6.0). Section was blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.