

## TRAPPC9 Ab

[References\(1\)](#) [Images\(3\)](#)

Cat.#: DF12489                      Concn.: ~1mg/ml                      Mol.Wt.: 128-140 kDa  
Size:                                      Source: Rabbit                          Clonality: Polyclonal

Application:                          WB 1:500-1:2000, IHC 1:50-1:200, IF/ICC 1:100-1:500  
\*The optimal dilutions should be determined by the end user.

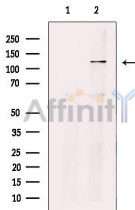
Reactivity:                            Human,Mouse,Rat,Zebrafish

Storage:                                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.

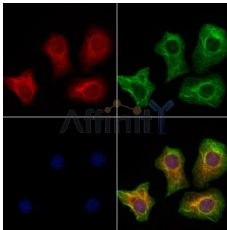
Purification:                         The antiserum was purified by peptide affinity chromatography using SulfoLink™ Coupling Resin (Thermo Fisher Scientific).

Immunogen:                          A synthesized peptide derived from human TRAPPC9, corresponding to a region within the internal amino acids.

Uniprot:                                Q96Q05



Western blot analysis of extracts from mouse brain tissue, using TRAPPC9 Ab. The lane on the left was treated with blocking peptide.



DF12489 staining Hela 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(DF12489 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. The nuclear counter stain is DAPI(blue).

**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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