[Primary Antibody]

phospho-DAB2 (Ser24) Rabbit pAb

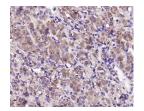


Pig, Cow,

www.bioss.com.cn sales@bioss.com.cn techsupport@bioss.com.cn 400-901-9800

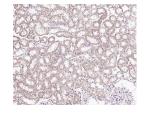
– DATASHEET –		400-901-9800
Host: Rabbit	Isotype: IgG	Applications: IHC-P (1:100-500) IHC-F (1:100-500)
Clonality: Polyclonal GenelD: 1601	SWISS: P98082	IF (1:100-500) Flow-Cyt (2ug/Test)
	4) gated synthesised phosphopeptide derived from humar nd the phosphorylation site of Ser24: AP(p-S)KK.	Reactivity: Human, Mouse (predicted: Rat, Pig, C Dog)
Purification: affinity purified by Protein A Concentration: 1mg/ml		Predicted MW.: ^{85 kDa}
Storage: 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol. Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles.		Subcellular Location: Cytoplasm
expressed i or absent fr tumor supp an adaptor (a guanine proline-rich pathways b Alternative	ncodes a mitogen-responsive phosphoprotein. It is n normal ovarian epithelial cells, but is down-regulated rom ovarian carcinoma cell lines, suggesting its role as pressor. This protein binds to the SH3 domains of GRB2 protein that couples tyrosine kinase receptors to SOS nucleotide exchange factor for Ras), via its C-terminal n sequences, and may thus modulate growth factor/Ras by competing with SOS for binding to GRB2. Ily spliced transcript variants encoding different ave been found for this gene. [provided by RefSeq, Oct	a ,

– VALIDATION IMAGES

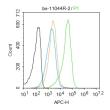


2011].

Paraformaldehyde-fixed, paraffin embedded (human adrenal gland); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Incubation with (phospho-DAB2 (Ser24)) Polyclonal Antibody, Unconjugated (bs-11044R) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructionsand DAB staining.



Paraformaldehyde-fixed, paraffin embedded (mouse kidney); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Incubation with (phospho-DAB2 (Ser24)) Polyclonal Antibody, Unconjugated (bs-11044R) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructionsand DAB staining.



Blank control: Mouse spleen. Primary Antibody (green line): Rabbit Anti-DAB2 antibody (bs-11044R) Dilution: 2µg /10^6 cells; Isotype Control Antibody (orange line): Rabbit IgG . Secondary Antibody : Goat anti-rabbit IgG-AF647 Dilution: $1\mu g$ /test. Protocol The cells were fixed with 4% PFA (10min at room temperature) and then permeabilized with 90% ice-cold methanol for 20 min at-20°C. The cells were then incubated in 5%BSA to block non-specific protein-protein interactions for 30 min at room temperature .Cells stained with Primary Antibody for 30 min at room temperature. The secondary antibody used for 40 min at room temperature. Acquisition of 20,000 events was performed.

- SELECTED CITATIONS -

• [IF=4.7] Gloria Riitano. et al. Role of Lipid Rafts on LRP8 Signaling Triggered by Anti-β2-GPI Antibodies in Endothelial Cells. BIOMEDICINES. 2023 Dec;11(12):3135 WB ;Human. 10.3390/biomedicines11123135