## [ Primary Antibody ]

## phospho-FAK (Tyr407) Rabbit pAb



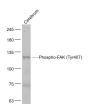
www.bioss.com.cn sales@bioss.com.cn techsupport@bioss.com.cn 400-901-9800

– DATASHEET –––––		400-901-9800
Host: Rabbit	Isotype: IgG	Applications: WB (1:500-2000)
Clonality: Polyclonal		IHC-P (1:100-500) IHC-F (1:100-500)
GenelD: 5747	SWISS: Q05397	<b>IF</b> (1:100-500)
Target: FAK (Tyr407)		Flow-Cyt (1µg /Test)
Immunogen: KLH conjugated Synthesised phosphopeptide derived from human FAK around the phosphorylation site of Tyr407: DT(p-Y)TM.		(predicted: Rat, Rabbit,
Purification: affinity purified by Protein A		Sheep, Cow, Dog, Horse)
Concentration: 1mg/ml		Predicted
<b>Storage:</b> 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol.		MW.: <sup>116</sup> kDa
, ,	re at -20°C for one year. Avoid repeated	Subcellular Cell membrane ,Cytoplasm Location: ,Nucleus
<b>Background:</b> Non-receptor protein-tyrosine kinase implicated in signaling pathways involved in cell motility, proliferation and apoptosis. Activated by tyrosine-phosphorylation in response to either integrin clustering induced by cell adhesion or antibody cross- linking, or via G-protein coupled receptor (GPCR) occupancy by ligands such as bombesin or lysophosphatidic acid, or via LDL receptor occupancy. Plays a potential role in oncogenic		is. ss- <sup>,</sup> by

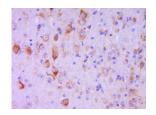
transformations resulting in increased kinase activity. [SUBCELLULAR LOCATION] Cell junction, focal adhesion. Cell membrane; Peripheral membrane protein; Cytoplasmic side.

Note=Constituent of focal adhesions.

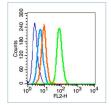
## – VALIDATION IMAGES



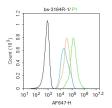
Sample: Cerebrum (Mouse) Lysate at 40 ug Primary: Anti- Phospho-FAK (Tyr407) (bs-3164R) at 1/1000 dilution Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution Predicted band size: 116 kD Observed band size: 116 kD



Paraformaldehyde-fixed, paraffin embedded (mouse brain tissue); 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; Antibody incubation with (P-FAK (Tyr407)) Polyclonal Antibody, Unconjugated (bs-3164R) at 1:400 overnight at 4°C, followed by a conjugated secondary (sp-0023) for 20 minutes and DAB staining.



Blank control (blue line): Hep G2 (fixed with 70% ethanol (Overmight at 4°C) and then permeabilized with 90% ice-cold methanol for 30 min at -20°C). Primary Antibody (green line): Rabbit Anti-Phospho-FAK (Tyr407) antibody (bs-3164R),Dilution: 1µg /10^6 cells; Isotype Control Antibody (orange line): Rabbit IgG . Secondary Antibody (white blue line): Goat antirabbit IgG-PE,Dilution: 1µg /test.



Blank control:A431. Primary Antibody (green line): Rabbit Anti-Phospho-FAK (Tyr407) antibody (bs-3164R) Dilution: 1µg /10^6 cells; Isotype Control Antibody (orange line): Rabbit IgG . Secondary Antibody : Goat anti-rabbit IgG-AF647 Dilution: 1µg /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 nonspecific 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=4.6] Giuseppe Prencipe. et al. Amphiregulin Orchestrates the Paracrine Immune-Suppressive Function of Amniotic-Derived Cells Through its Interplay with COX-2/PGE2/EP4 Axis. ISCIENCE. 2024 七月 13 WB ;Human. 39156643
- [IF=4.6] Giuseppe Prencipe. et al.Amphiregulin orchestrates the paracrine immune-suppressive function of amnioticderived cells through its interplay with COX-2/PGE<sub>2</sub>/EP4 axis.iscience.2024 Jul 14;27(8):110508. Western blot ;Sheep. 39156643