bsm-60586R

[Primary Antibody]

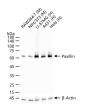
Paxillin Recombinant Rabbit mAb



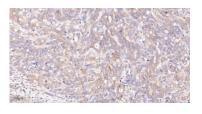
www.bioss.com.cn sales@bioss.com.cn techsupport@bioss.com.cn 400-901-9800

- DATASHEET		400-901-9800
Host: Rabbit	Isotype: IgG	Applications: WB (1:500-2000)
Clonality: Recombinant	CloneNo.: E2E8	IHC-P (1:50-200) IHC-F (1:50-200)
Target: Paxillin Purification: affinity purified by Protein A		IF (1:50-200) Flow-Cyt (1:50-100) ICC/IF (1:50-200)
Concentration: 1mg/ml		Reactivity: Human, Mouse, Rat
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
cell adhesion and migra variety of processes incl wound repair, inflamma motifs, LIM domains, SH docking sites for cytosk Pyk 2, Src), serine/threo	skeletal adapter protein involved in on of focal adhesions, which are critic tion. This in turn plays a role in a wid luding embryogenesis, organogenesis ation and cancer. Paxillin contains LD I3 and SH2 binding domains that serv eletal proteins, tyrosine kinases (e.g., nine kinases, GTPase activating prote eins (e.g., Actin, Vinculin, Crk).	e s, /e as FAK,

- VALIDATION IMAGES -



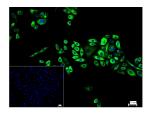
25 ug total protein per lane of various lysates (see on figure) probed with Paxillin monoclonal antibody, unconjugated (bsm-60586R) at 1:1000 dilution and 4°C overnight incubation. Followed by conjugated secondary antibody incubation at r.t. for 60 min.



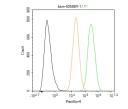
Paraformaldehyde-fixed, paraffin embedded Human Ovarian Cancer; Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15 min; The section was incubated with Paxillin Monoclonal Antibody, Unconjugated (bsm-60586R) at 1:200 overnight at 4°C, followed by conjugation to the bs-0295G-HRP and DAB (C-0010) staining.



Paraformaldehyde-fixed, paraffin embedded Rat Cerebrum; Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15 min; The section was incubated with Paxillin Monoclonal Antibody, Unconjugated (bsm-60586R) at 1:200 overnight at 4°C, followed by conjugation to the bs-0295G-HRP and DAB (C-0010) staining.



4% Paraformaldehyde-fixed PC-3 (H) cell; Triton X-100 at r.t. for 20 min; Antibody incubation with (Paxillin) monoclonal Antibody, unconjugated (bsm-60586R) 1:100, 90 min at 37°C; followed by conjugated Goat Anti-Rabbit IgG antibody (green, bs-60295G-BF488) at 37°C for 90 min, DAPI (blue, C02-04002) was used to stain the cell nuclei. PBS instead of the primary antibody was used as the blank control.



The PC-3 (H) cells were fixed with 4% PFA (10 min at r.t.) and then permeabilized with 90% icecold methanol for 20 min at -20°C,the cells then were incubated in 5%BSA to block non-specific protein-protein interactions (30 min at r.t.), followed by secondary antibody incubation for 40 min at room temperature.Primary Antibody (green):Rabbit Anti-Paxillin antibody (bsm-60586R,1:100); Isotype Control (orange): Rabbit IgG (bs-0295P). Blank control (black): PBS. Acquisition of 20,000 events was performed.