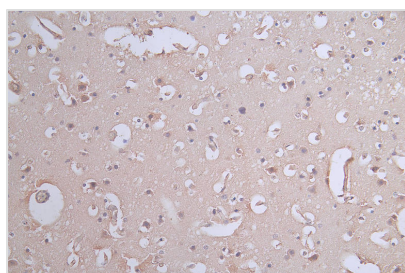




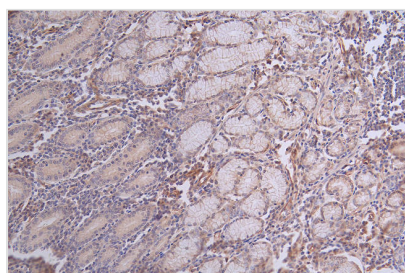
# PXN Recombinant Monoclonal Antibody

<b>Product Code</b>	CSB-RA017450A0HU
<b>Storage</b>	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.
<b>Uniprot No.</b>	P49023
<b>Immunogen</b>	A synthesized peptide derived from Human PXN
<b>Species Reactivity</b>	Human
<b>Tested Applications</b>	ELISA, IHC, FC; Recommended dilution: IHC:1:50-1:200, FC:1:50-1:200
<b>Form</b>	Liquid
<b>Conjugate</b>	Non-conjugated
<b>Storage Buffer</b>	Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol.
<b>Purification Method</b>	Affinity-chromatography
<b>Isotype</b>	Rabbit IgG
<b>Clonality</b>	Monoclonal
<b>Product Type</b>	Recombinant Antibody
<b>Immunogen Species</b>	Homo sapiens (Human)
<b>Research Area</b>	Cancer;Signal transduction
<b>Gene Names</b>	PXN
<b>Clone No.</b>	32E1

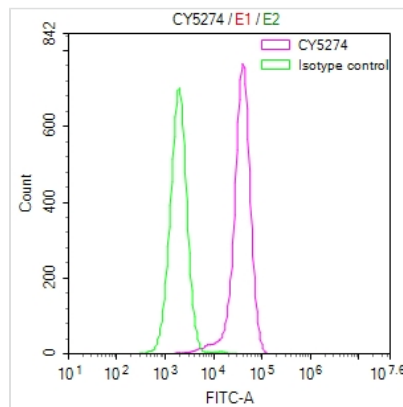
## Image



IHC image of CSB-RA017450A0HU diluted at 1:50 and staining in paraffin-embedded human brain tissue performed on a Leica Bond<sup>TM</sup> 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 Goat anti-rabbit polymer IgG labeled by HRP and visualized using 0.62% DAB.



IHC image of CSB-RA017450A0HU diluted at 1:50 and staining in paraffin-embedded human breast cancer performed on a Leica Bond<sup>TM</sup> 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 Goat anti-rabbit polymer IgG labeled by HRP and visualized using 0.62% DAB.



Overlay Peak curve showing HepG2 cells stained with CSB-RA017450A0HU (red line) at 1:50. The cells were fixed in 4% formaldehyde and permeated by 0.2% TritonX-100. Then 10% normal goat serum to block non-specific protein-protein interactions followed by the antibody ( $1\mu\text{g}/1 \times 10^6$  cells) for 45min at 4?. The secondary antibody used was FITC-conjugated Goat Anti-rabbit IgG(H+L) at 1:200 dilution for 35min at 4?. Control antibody (green line) was rabbit IgG ( $1\mu\text{g}/1 \times 10^6$  cells) used under the same conditions. Acquisition of >10,000 events was performed.

## Description

The production of the PXN recombinant monoclonal antibody relies on in vitro expression systems developed through the cloning of PXN antibody DNA sequences sourced from immunoreactive rabbits. The immunogen employed is a synthesized peptide derived from human paxillin. Subsequently, the PXN antibody genes are inserted into plasmid vectors, and these recombinant plasmid vectors are transfected into host cells to facilitate antibody expression. Following expression, the PXN recombinant monoclonal antibody is subjected to affinity-chromatography purification. It undergoes comprehensive testing in ELISA, IHC, and FC applications, confirming its reactivity with the human PXN protein.

Paxillin (PXN) is a versatile protein that plays a central role in cell adhesion, signaling, and cytoskeletal organization. Its functions are critical for various cellular processes, including cell adhesion, migration, and tissue development. Dysregulation of paxillin can have significant implications in diseases like cancer and neurological disorders.