

beta Catenin (phospho Ser33) antibody [HL5105]

Cat. No. GTX645299

Host	Rabbit
Clonality	Monoclonal
Isotype	IgG
Applications	WB
Reactivity	Human, Mouse, Rat

Package
100 µl, 25 µl

Applications

Application Note

*Optimal dilutions/concentrations should be determined by the researcher.

Suggested dilution	Recommended dilution
WB	1:500-1:3000

Not tested in other applications.

Observed MW (kDa) 97 kDa.

Properties

Form	Liquid
Buffer	PBS
Preservative	No preservatives
Storage	Store as concentrated solution. Centrifuge briefly prior to opening vial. For short-term storage (1-2 weeks), store at 4°C. For long-term storage, aliquot and store at -20°C or below. Avoid multiple freeze-thaw cycles.
Concentration	1 mg/ml (Please refer to the vial label for the specific concentration.)
Immunogen	Carrier-protein conjugated synthetic peptide surrounding phospho Ser33 of human beta Catenin. The exact sequence is proprietary.
Purification	Affinity purified by Protein A.
Conjugation	Unconjugated

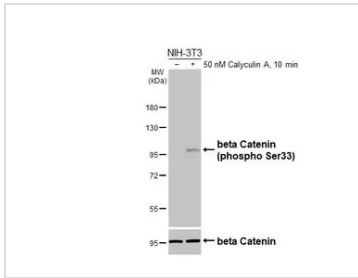
Note

For laboratory research use only. Not for any clinical, therapeutic, or diagnostic use in humans or animals. Not for animal or human consumption.

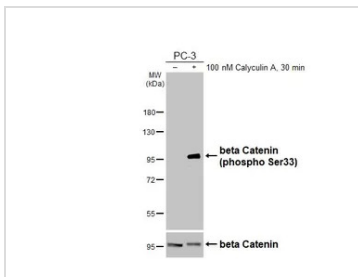
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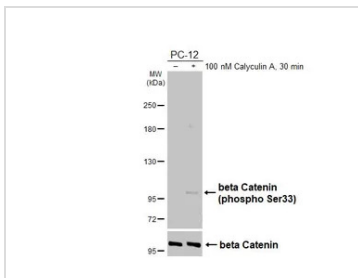
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DATA IMAGES

GTX645299 WB Image

Untreated (–) and treated (+) NIH-3T3 whole cell extracts (30 µg) were separated by 7.5% SDS-PAGE, and the membranes were blotted with beta Catenin (phospho Ser33) antibody [HL5105] (GTX645299) diluted at 1:1000 and beta Catenin antibody [HL1753] (GTX637403) diluted at 1:1000. The HRP-conjugated anti-rabbit IgG antibody (GTX213110-01) was used to detect the primary antibody, and the signal was developed with Trident femto Western HRP Substrate.


GTX645299 WB Image

Untreated (–) and treated (+) PC-3 whole cell extracts (30 µg) were separated by 7.5% SDS-PAGE, and the membranes were blotted with beta Catenin (phospho Ser33) antibody [HL5105] (GTX645299) diluted at 1:1000 and beta Catenin antibody [HL1753] (GTX637403) diluted at 1:1000. The HRP-conjugated anti-rabbit IgG antibody (GTX213110-01) was used to detect the primary antibody, and the signal was developed with Trident femto Western HRP Substrate.


GTX645299 WB Image

Untreated (–) and treated (+) PC-12 whole cell extracts (30 µg) were separated by 5% SDS-PAGE, and the membrane was blotted with beta Catenin (phospho Ser33) antibody [HL5105] (GTX645299) diluted at 1:500 and beta Catenin antibody [HL1753] (GTX637403) diluted at 1:500. The HRP-conjugated anti-rabbit IgG antibody (GTX213110-01) was used to detect the primary antibody.



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