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Research Use Only. Not for diagnostic or therapeutic use.

EB11651 - Goat Anti-PCNA (aa111-122) Antibody

Size: 100µg specific antibody in 200µl



Target Protein

Principal Names: cyclin, DNA polymerase delta auxiliary protein, MGC8367, OTTHUMP0000030189, OTTHUMP0000030190, PCNA, proliferating cell nuclear

antigen

Official Symbol: PCNA

Accession Number(s): NP_002583.1

Human GeneID(s): 5111

Non-Human GenelD(s): 18538 (mouse), 25737 (rat)

Important Comments: Reported variants represent identical protein: NP_872590.1,

NP_002583.1

Immunogen

Peptide with sequence C-EAPNQEKVSDYE, from the internal region of the protein sequence according to NP_002583.1.

Please note the peptide is available for sale.

Purification and Storage

Purified from goat serum by ammonium sulphate precipitation followed by antigen affinity chromatography using the immunizing peptide.

Supplied at 0.5 mg/ml in Tris saline, 0.02% sodium azide, pH7.3 with 0.5% bovine serum albumin.

Aliquot and store at -20°C. Minimize freezing and thawing.

Applications Tested

Peptide ELISA: antibody detection limit dilution 1:128000.

Western blot: Approx 30kDa band observed in lysates of cell line Jurkat (calculated MW of 28.8kDa according to Human NP_002583.1). Approx.35kDa band was observed in lysates of cell line NIH3T3 and in Mouse and Rat Testis, and approx 33kDa in Pig Spleen lysates. (calculated MW of 28.8kDa according to Mouse NP_035175.1, Rat NP_071776.1 and Pig NP_001278854.1). The observed molecular weight corresponds to earlier findings in literature with different antibodies (Gilljam et al, , J Cell Biol. 2009 Sep 7;186(5):645-54.). Recommended concentration: 0.03-0.1μg/ml.

IHC: Paraffin embedded Human Tonsil. Recommended concentration: 2.5µg/ml.

Additional validation: This antibody has been successfully used in the following paper: Sikorski et al. (2018) PMID: 30377371.

Species Reactivity

Tested: Human, Mouse, Rat, Pig

Expected from sequence similarity: Human, Mouse, Rat, Dog, Pig, Cow

Specific Reference

This antibody has been successfully used in the following paper:

Krzysztof Sikorski, Adi Mehta, Marit Inngjerdingen, Flourina Thakor, Simon Kling, Tomas Kalina, Tuula A. Nyman, Maria Ekman Stensland, Wei Zhou, Gustavo A. De Souza, Lars Holden, Jan Stuchly, Markus Templin and Fridtjof Lund-Johansen

A high-throughput pipeline for validation of antibodies

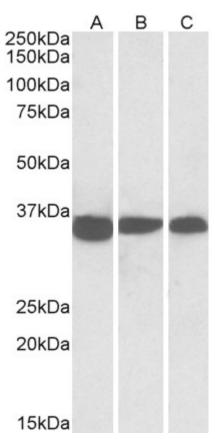
Nat Methods. 2018 Nov;15(11):909-912

PMID: 30377371



EB11651 (0.05µg/ml) staining of Jurkat lysate (35µg protein in RIPA buffer). Primary incubation was 1 hour.

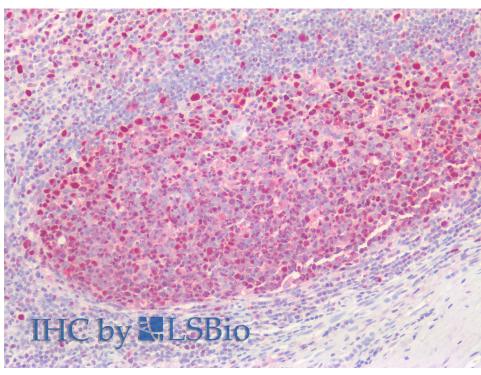
Detected by chemiluminescence.



EB11650 (0.03μg/ml) staining of NIH3T3 (A), Mouse Testis (B) and Rat Testis (C) lysates (35μg protein in RIPA buffer). Primary incubation was 1 hour. Detected by chemiluminescence.

250kDa 150kDa 100kDa 75kDa 50kDa 37kDa 25kDa 20kDa

EB11651 (0.3 μ g/ml) staining of Pig Spleen lysate (35 μ g protein in RIPA buffer). Primary incubation was 1 hour. Detected by chemiluminescence.



EB11651 (2.5μg/ml) staining of paraffin embedded Human Tonsil. Steamed antigen retrieval with citrate buffer pH 6, AP-staining.