



## International Office

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**Research Use Only. Not for  
diagnostic or therapeutic use.**

## EB08932 - Goat Anti-SEPT6 Antibody

Size: 100µg specific antibody in 200µl



### Target Protein

**Principal Names:** SEPT6, septin 6, KIAA0128, MGC16619, MGC20339, SEP2, SEPT2, septin 2

**Official Symbol:** SEPTIN6

**Accession Number(s):** NP\_665798.1; NP\_055944.2; NP\_665801.1

**Human GeneID(s):** [23157](#)

**Important Comments:** This antibody is expected to recognize the reported isoforms NP\_665798.1, NP\_055944.2 and NP\_665801.1. Reported variants NP\_665799.1 and NP\_665798.1 represent identical protein.

### Immunogen

Peptide with sequence C-DEVNAFKQRKTA, from the internal region of the protein sequence according to NP\_665798.1; NP\_055944.2; NP\_665801.1.

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 1:4000.

**Western blot:** Approx 50kDa band observed in Human Testis, Duodenum, Kidney and Tonsil lysates, and in lysates of cell lines Daudi, Jurkat and MOLT-4 (calculated MW of 49.7kDa according to NP\_055944.2). Recommended concentration: 0.1-1µg/ml. Primary incubation 1 hour at room temperature.

**IHC:** In paraffin embedded Human Tonsil shows specific staining in dividing cells just outside a germinal centre. Recommended concentration: 3-5µg/ml.

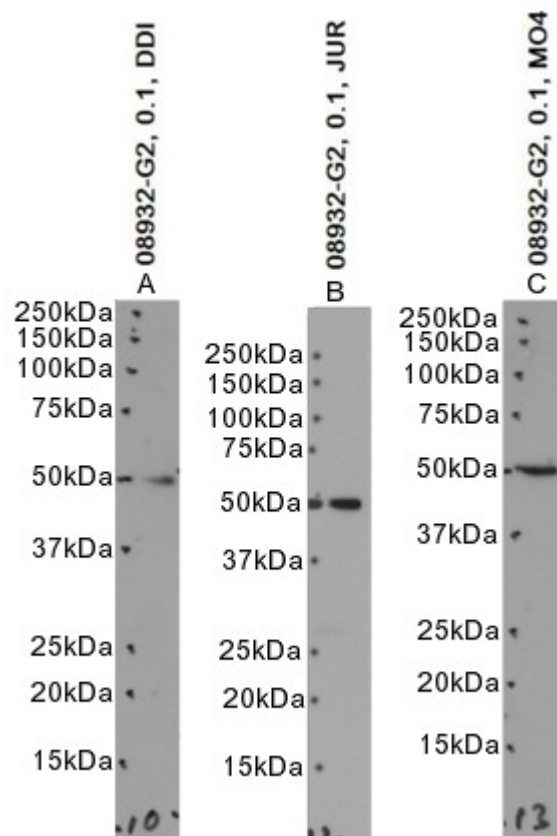
**Immunofluorescence:** Strong expression of the protein seen in A431 and U2OS cells. Recommended concentration: 10µg/ml.

**Flow Cytometry:** Flow cytometric analysis of A431 cells. Recommended concentration: 10ug/ml.

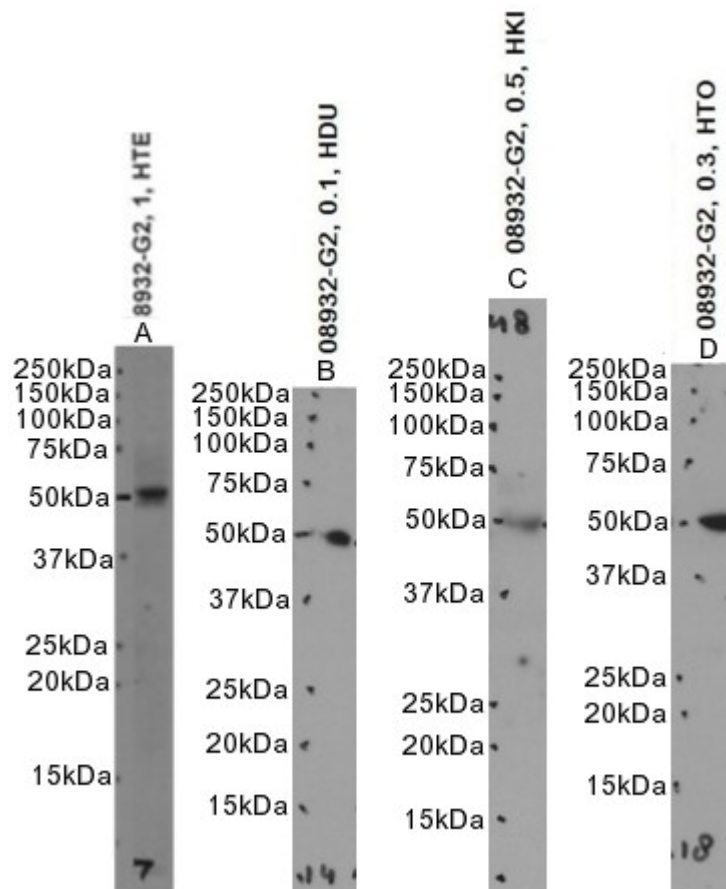
### Species Reactivity

**Tested:** Human

**Expected from sequence similarity:** Human, Cow

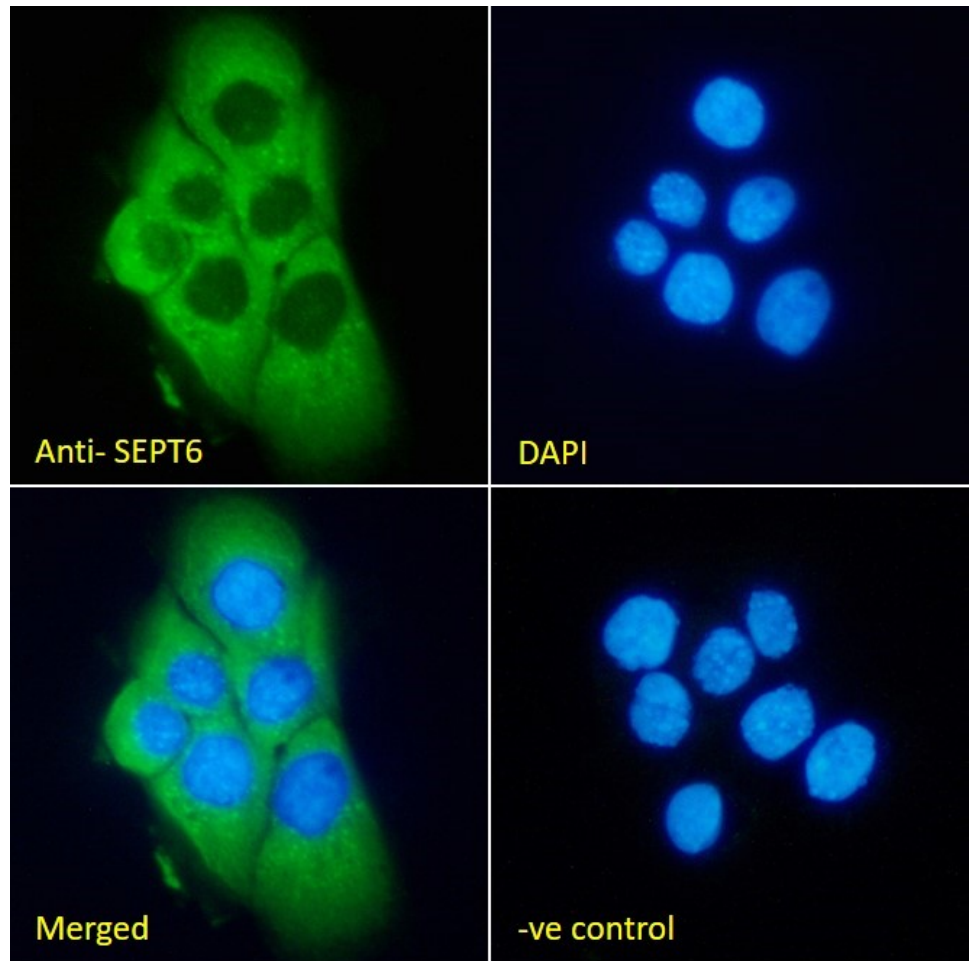


EB08932 optimised QC. Primary incubation 1 hour at room temperature.  
 Images A, B, C: Daudi, Jurkat, MOLT-4 cell lysates at primary Ab concentration 0.1ug/ml  
 (Loaded 35µg protein in RIPA buffer, per lane). Detected by chemiluminescence.

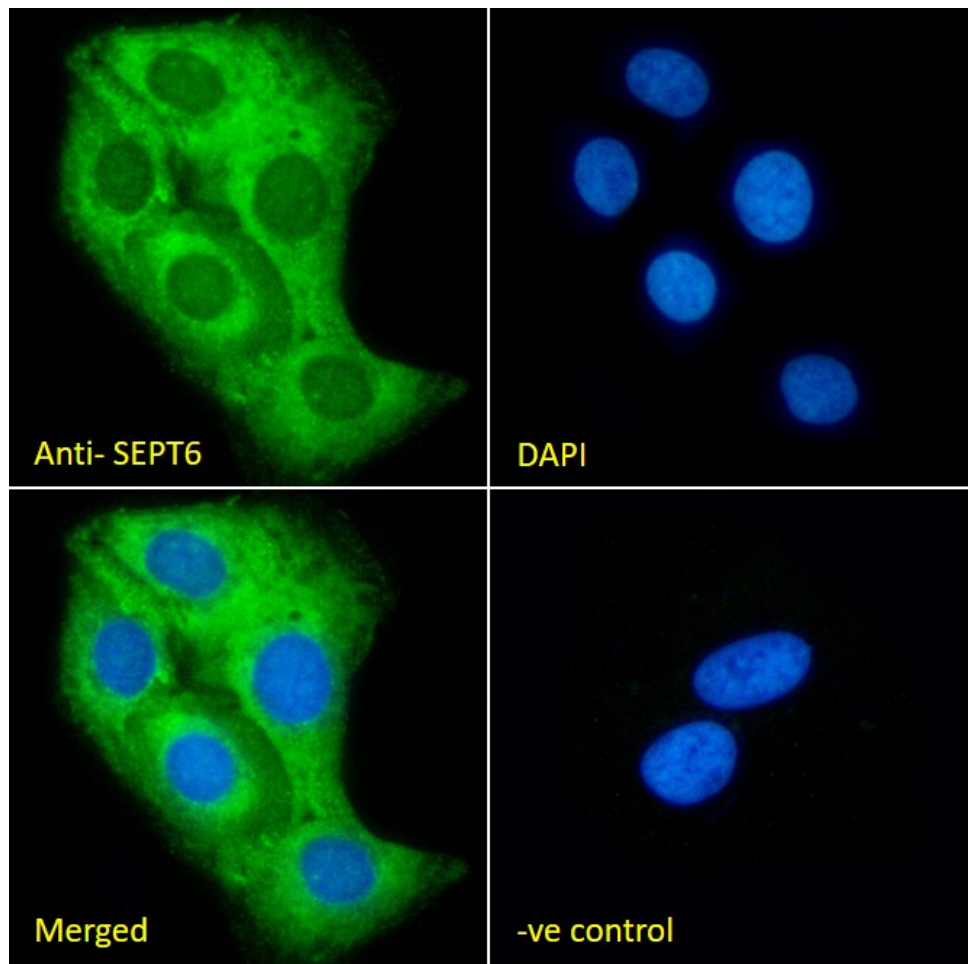


EB08932 optimised QC. Primary incubation 1 hour at room temperature.

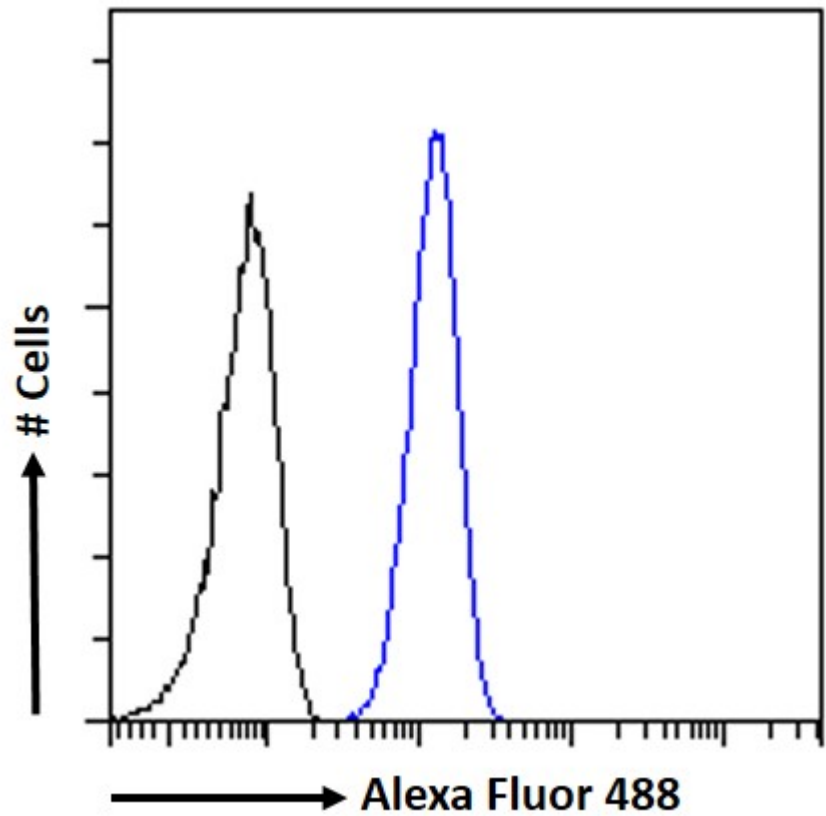
Image A: Human Testes lysate at primary Ab concentration 1ug/ml, Image B: Human Duodenum lysate at primary Ab concentration 0.1ug/ml, Image C: Human Kidney lysate at primary Ab concentration 0.5ug/ml, Image D: Human Tonsil lysate at primary Ab concentration 0.3ug/ml (Loaded 35µg protein in RIPA buffer, per lane). Detected by chemiluminescence.



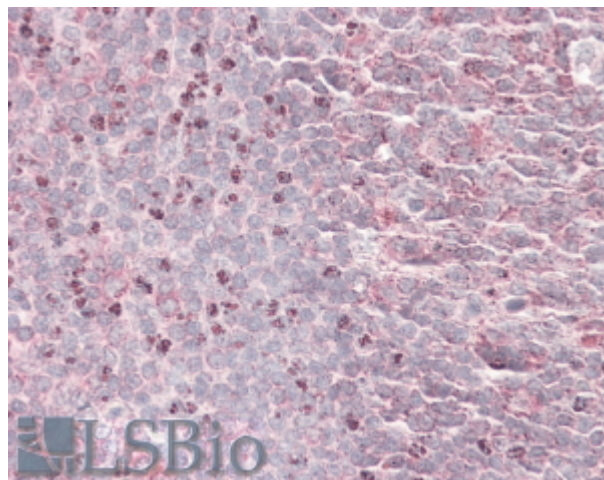
EB08932 Immunofluorescence analysis of paraformaldehyde fixed A431 cells, permeabilized with 0.15% Triton. Primary incubation 1hr (10ug/ml) followed by Alexa Fluor 488 secondary antibody (2ug/ml), showing cytoplasmic staining. The nuclear stain is DAPI (blue). Negative control: Unimmunized goat IgG (10ug/ml) followed by Alexa Fluor 488 secondary antibody (2ug/ml).



EB08932 Immunofluorescence analysis of paraformaldehyde fixed U2OS cells, permeabilized with 0.15% Triton. Primary incubation 1hr (10ug/ml) followed by Alexa Fluor 488 secondary antibody (2ug/ml), showing cytoplasmic staining. The nuclear stain is DAPI (blue). Negative control: Unimmunized goat IgG (10ug/ml) followed by Alexa Fluor 488 secondary antibody (2ug/ml).



EB08932 Flow cytometric analysis of paraformaldehyde fixed A431 cells (blue line), permeabilized with 0.5% Triton. Primary incubation 1hr (10ug/ml) followed by Alexa Fluor 488 secondary antibody (1ug/ml). IgG control: Unimmunized goat IgG (black line) followed by Alexa Fluor 488 secondary antibody.



EB08932 (3.8µg/ml) staining of paraffin embedded Human Tonsil. Steamed antigen retrieval with citrate buffer pH 6, AP-staining.