

## UK Office

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

## EB11207 - Goat Anti-Vimentin Antibody

Size: 100µg specific antibody in 200µl



### Target Protein

**Principal Names:** FLJ36605, vimentin, VIM

**Official Symbol:** VIM

**Accession Number(s):** NP\_003371.2

**Human GeneID(s):** [7431](#)

**Non-Human GeneID(s):** 22352 (mouse), 81818 (rat)

### Immunogen

Peptide with sequence C-QVINETSQHDDLE, from the C Terminus of the protein sequence according to NP\_003371.2.

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 55kDa band observed in lysates of cell line HeLa and Jurkat and in Mouse Ovary lysates, and approx. 55-60kDa band in Rat Ovary lysates (calculated MW of 53.7kDa according to Human NP\_003371.2, Mouse NP\_035831.2 and Rat NP\_112402.1). Recommended concentration: 0.1-2µg/ml.

**Flow Cytometry:** Flow cytometric analysis of HeLa cells. Recommended concentration: 10ug/ml. **Immunofluorescence:** Strong expression of the protein seen in the cytoplasm/Intermediate filaments of U2OS cells. Recommended concentration: 5µg/ml.

### Species Reactivity

**Tested:** Human, Mouse, Rat

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

EB11207 (2µg/ml) staining of HeLa (A) and Jurkat (B) lysates (35µg protein in RIPA buffer). Primary incubation was 1 hour. Detected by chemiluminescence.

EB11207 (0.1µg/ml) staining of Mouse (A) and (2µg/ml) Rat (B) Ovary lysate (35µg protein in RIPA buffer). Primary incubation was 1 hour. Detected by chemiluminescence.

Flow cytometric analysis of paraformaldehyde fixed HeLa cells (blue line), permeabilized with 0.5% Triton. Primary incubation 1hr (10ug/ml) followed by Alexa Fluor 488 secondary antibody (0.4ug/ml). IgG control: Unimmunized goat IgG (black line) followed by Alexa Fluor 488 secondary antibody.

EB11207 Immunofluorescence analysis of paraformaldehyde fixed U2OS cells, permeabilized with 0.15% Triton. Primary incubation 1hr (5ug/ml) followed by Alexa Fluor 488 secondary antibody (2ug/ml), showing cytoplasmic/Intermediate filament staining. The nuclear stain is DAPI (blue). Negative control: Unimmunized goat IgG (5ug/ml) followed by Alexa Fluor 488 secondary antibody (2ug/ml).