



## UK Office

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

## EB05545 - Goat Anti-ARL2 Antibody

Size: 100µg specific antibody in 200µl



### Target Protein

**Principal Names:** ARL2, ADP-ribosylation factor-like 2, ARFL2, ADP=ribosylation factor-like 2

**Official Symbol:** ARL2

**Accession Number(s):** NP\_001658.2

**Human GeneID(s):** [402](#)

### Immunogen

Peptide with sequence C-LDDISSRIFTAD, from the C Terminus of the protein sequence according to NP\_001658.2.

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 1:64000.

**Western blot:** Approx. 18kDa band observed in lysates of cell line A431 and approx. 20kDa in nuclear lysates of cell line HeLa (calculated MW of 20.9kDa according to NP\_001658.2). Recommended concentration: 0.5-1µg/ml. Primary incubation 1 hour at room temperature.

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

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

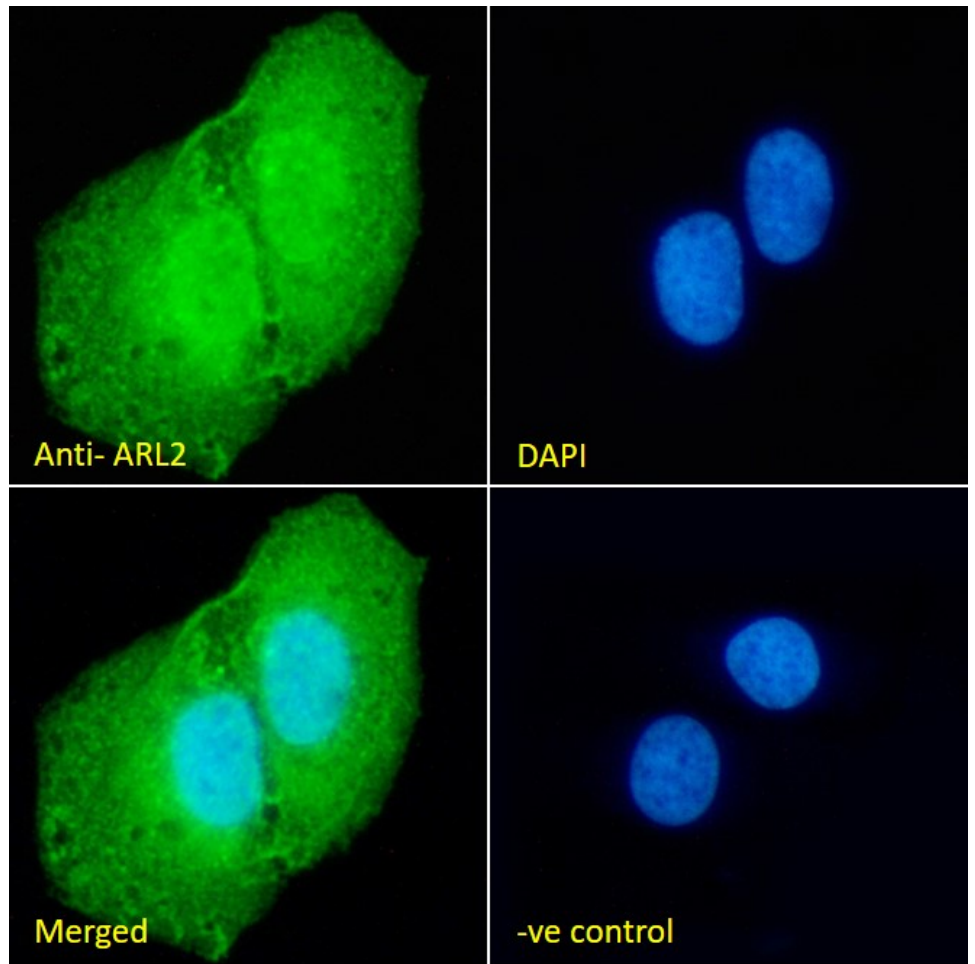
### Species Reactivity

**Tested:** Human

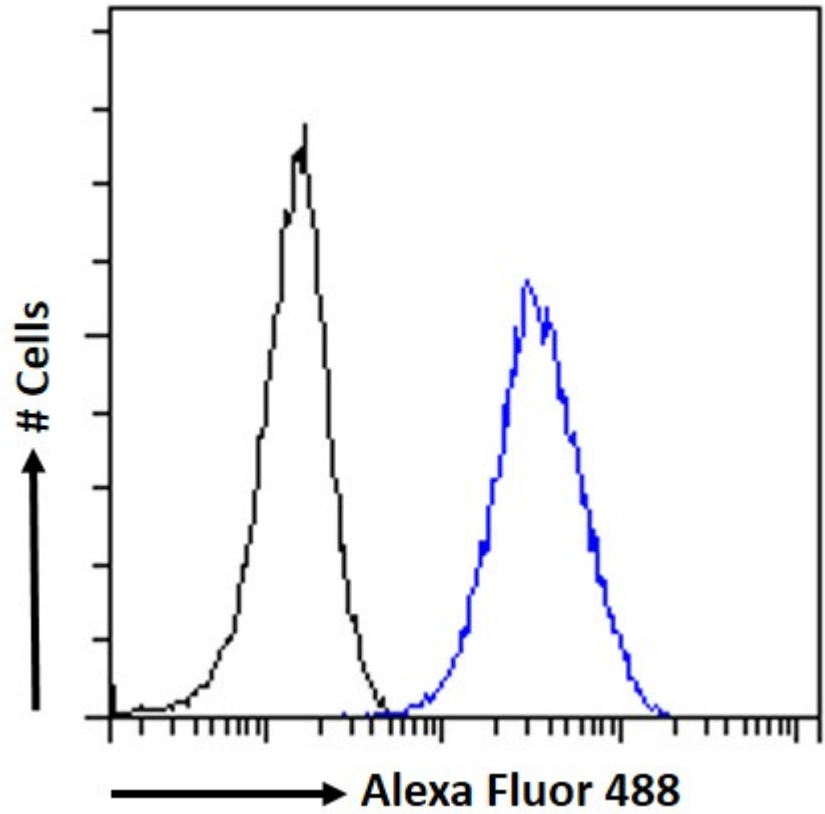
**Expected from sequence similarity:** Human, Dog



EB05545 (0.5 $\mu$ g/ml) staining of A431 (A) and (1 $\mu$ g/ml) HeLa nuclear (B) cell lysate (30 $\mu$ g protein in RIPA buffer).  
Detected by chemiluminescence.



EB05545 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 nuclear and cytoplasmic staining. The nuclear stain is DAPI (blue). Negative control: Unimmunized goat IgG (10ug/ml) followed by Alexa Fluor 488 secondary antibody (2ug/ml).



EB05545 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.